

091965594

GenCore version 5.1.6
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OM protein - protein search, using sw mode:

Run on: September 27, 2003, 06:45:13 : Search time 23 Seconds

(without alignments)
372.124 Million cell updates/sec

Title: US-09-965-594-1

Perfect score: 953

Sequence: 1 MAPITAYAQTKGLLCIIT.....GVAKAVDFIPVESLETTMRS 132

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 127863 seqs, 47026705 residues

Total number of hits satisfying chosen parameters: 127863

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 99%

Listing first 45 summaries

Database : SwissProt_41.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result NO.	Score	Query Match %	Length	DB ID	Description
1	943	99.0	3011	1	POLG_HCV1
2	927	97.3	3011	1	POLG_HCVH
3	892	93.5	3010	1	POLG_HCVTW
4	891	93.5	3010	1	POLG_HCVBK
5	887	93.1	3010	1	POLG_HCVJA
6	884	92.8	3010	1	POLG_HCVJT
7	714	74.9	3033	1	POLG_HCVJ9
8	712	74.7	3033	1	POLG_HCVJ6
9	87	9.1	209	1	PAAD_PSEAE
10	84	8.8	321	1	HHOA_ARATH
11	82	8.6	452	1	AAMP_HUMAN
12	82	8.6	485	1	Y136_TREPA
13	80.5	8.4	437	1	DEGL_ARATH
14	75	7.9	253	1	CAC3_BOVIN
15	74.5	7.8	415	1	ZP3_RABIT
16	74.5	7.8	776	1	HYPF_AZOFI
17	74.5	7.8	911	1	TB11_RHIMB
18	74	7.8	326	1	PANE_RHILQ
19	73.5	7.7	263	1	GRAK_MOUSE
20	73	7.7	730	1	HELS_METVA
21	72.5	7.6	257	1	GRAM_HUMAN
22	72	7.6	627	1	SAD1_MOUSE
23	72	7.6	1527	1	CAIH_MOUSE
24	72	7.6	2663	1	GENE_HUMAN
25	72	7.6	3491	1	ERVL_SACER
26	71.5	7.5	248	1	GRAD_MOUSE
27	71.5	7.5	323	1	VPRT_SRRVH
28	71	7.5	219	1	SPRI_IPOBA
29	71	7.5	336	1	UL16_EBV
30	71	7.5	529	1	PLG2_RALSO
31	71	7.5	1180	1	ITAI_RAT
32	70.5	7.4	264	1	CTSL_HUMAN
33	70.5	7.4	659	1	VST2_HEVME

RESULT 1

ID	POLG_HCV1	STANDARD;	PRT;	3011 AA.
AC	P26664;			
DT	01-AUG-1992	(Rel. 23, Created)		
DT	01-AUG-1992	(Rel. 23, Last sequence update)		
DT	15-SEP-2003	(Rel. 42, Last annotation update)		
DE	Genome polyprotein [Contains: Capsid protein C (Core protein) (P22); Envelope glycoprotein E1 (GP32) (GP35); Envelope glycoprotein E2 (GP68) (GP70) (NS1); Protein P7: Nonstructural protein NS2 (P21) (EC 3.4.22.-); Protease/helicase NS3 (P70) (Hepacivirin) (EC 3.4.21.96); Nonstructural protein NS4A (P4); Nonstructural protein NS4B (P27); Nonstructural protein NS5A (P56); Nonstructural protein NS5B (P66) (P70) (RNA-directed RNA polymerase) (EC 2.7.7.48)].			
DE	Hepatitis C virus (isolate 1) (HCV).			
OS	Hepatitis C virus positive-strand viruses, no DNA stage; Flaviviridae; Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae; Hepacivirus.			
OC	Hepacivirus.			
OX	NCBI-TaxID:11104;			
FN	SEQUENCE FROM N.A.			
RP	MEDLINE-91172826; PubMed-1848704;			
RA	Choo Q.-L., Richman K.H., Han J.H., Berger K., Lee C., Dong C., Gallegos C., Coit D., Medina-Selby A., Barr P.J., Weiner A.J., Bradley D.W., Kuo G., Houghton M.; "Genetic organization and diversity of the hepatitis C virus."; Proc. Natl. Acad. Sci. U.S.A. 88:2451-2455(1991).			
CC	!- FUNCTION: THE SMALL PROTEINS NS2A, NS2B, NS4A AND NS4B ARE HYDROPHOBIC, SUGGESTING A POSSIBLE MEMBRANE-RELATED FUNCTION. NS3 AND NS5 MAY PLAY A ROLE IN THE VIRAL RNA REPLICATION.			
CC	!- CATALYTIC ACTIVITY: Hydrolysis of four peptide bonds in the viral precursor polyprotein, commonly with Asp or Glu in the P6 position, Cys or Thr in P1 and Ser or Ala in P1'.			
CC	!- CATALYTIC ACTIVITY: N nucleoside triphosphate - N diphosphate + (RNA)(N).			
CC	!- SUBUNIT: THE VIRION OF THIS VIRUS IS A NUCLEOCAPSID COVERED BY A LIPOPROTEIN ENVELOPE. THE ENVELOPE CONSISTS OF TWO PROTEINS: PROTEIN M AND GLYCOPROTEIN E. THE NUCLEOCAPSID IS A COMPLEX OF PROTEIN C AND RNA.			
CC	!- SIMILARITY: THE PROTEASE BELONGS TO PEPTIDASE FAMILY S29.			
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CC	EMBL: M62321; AAA5676.1;			
DR	PIR: A39166; GNMVC3.			
DR	PDB: 1ALV; 16-FEB-99.			
DR	PDB: 1HEI; 25-NOV-98.			
DR	MEROPS: S29.001;			
DR	MEROPS: 039.001;			
DR	InterPro: IPR001410; DEAD.			
DR	InterPro: IPR002522; HCV_capsid.			

InterPro: IPR002521; HCV_core.
 DR InterPro: IPR002519; HCV_env.
 DR InterPro: IPR002531; HCV_NS1.
 DR InterPro: IPR002518; HCV_NS2.
 DR InterPro: IPR004109; HCV_NS3.
 DR InterPro: IPR006745; HCV_NS4.
 DR InterPro: IPR001490; HCV_NS4b.
 DR InterPro: IPR002868; HCV_NS5a.
 DR InterPro: IPR002166; HCV_RdRp.
 DR InterPro: IPR001650; Helicase_C.
 DR InterPro: IPR007095; RNA_pol_DS_PS.
 DR InterPro: IPR007094; RNA_pol_PSVir.
 DR Pfam: PF01543; HCV_capsid; 1.
 DR Pfam: PF01542; HCV_core; 1.
 DR Pfam: PF01539; HCV_env; 1.
 DR Pfam: PF01560; HCV_NS1; 1.
 DR Pfam: PF01538; HCV_NS2; 1.
 DR Pfam: PF01537; HCV_NS3; 1.
 DR Pfam: PF01006; HCV_NS4a; 1.
 DR Pfam: PF01001; HCV_NS4b; 1.
 DR Pfam: PF01506; HCV_NS5a; 1.
 DR Pfam: PF00271; helicase_C; 1.
 DR Pfam: PF00398; Viral_RdRp; 1.
 DR ProDom: PD186062; HCV_NS1; 1.
 DR SMART: SM00487; DEXDC; 1.
 KW Polyprotein; Glycoprotein; Transferase; RNA-directed RNA polymerase;
 KW Core protein; Coat protein; Envelope protein; Helicase; ATP-binding;
 KW Transmembrane; Nonstructural protein; Hydrolase; Serine protease;
 KW 3D-structure.
 FT INIT_MET 1 1 REMOVED FROM CAPSID PROTEIN C BY THE
 CELLULAR AMINOPEPTIDASE.
 FT CHAIN 1 115
 FT CHAIN 116 191
 FT CHAIN 192 383
 FT CHAIN 384 729
 FT CHAIN 730 1006
 FT CHAIN 1007 1515
 FT CHAIN 1516 1862
 FT CHAIN 1863 2013
 FT CHAIN 2014 3011
 FT CHAIN 3012 369
 FT TRANSMEM
 FT ACT_SITE 1083 1083
 FT ACT_SITE 1137 1137
 FT ACT_SITE 1165 1165
 FT ACT_SITE 1230 1237
 FT NP_BIND 1316 1319
 FT SITE
 FT CARBOHYD 196 196
 FT CARBOHYD 209 209
 FT CARBOHYD 234 234
 FT CARBOHYD 305 305
 FT CARBOHYD 417 417
 FT CARBOHYD 423 423
 FT CARBOHYD 430 430
 FT CARBOHYD 448 448
 FT CARBOHYD 476 476
 FT CARBOHYD 532 532
 FT CARBOHYD 540 540
 FT CARBOHYD 556 556
 FT CARBOHYD 576 576
 FT CARBOHYD 623 623
 FT CARBOHYD 645 645
 FT CARBOHYD 2041 2041
 FT CARBOHYD 2077 2077
 FT CARBOHYD 2240 2240
 FT CARBOHYD 2364 2364
 FT CARBOHYD 2789 2789
 SQ SEQUENCE 3011 AA; 327197 MW; 63F8C9447FCE5AF9 CRC64;
 Query Match 99.0%; Score 943; Db 1; Length 3011;
 Best Local Similarity 98.4%; Pred. No. 3.4e-82;
 Matches 179; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

Db 1026 LAPITAYAQQTGRLGGLCIITSLTGRKNOVEQVIVSTAAQTFLATCINGVCWTYVHCA 1085
 QY 61 GTRIATSPKGPVIQMYTNYVDKOLVGWPAPOGSKSLPTCGSSDIYIVTRHADVIPVRRR 120
 Db 1086 GTTRIATSPKGPVIQMYTNYVDKOLVGWPAPOGSKSLPTCGSSDIYIVTRHADVIPVRRR 1145
 QY 121 GDSRGLSLSPRISYLVKSGGGPGLCPAGHAGIFRAAVCTIRGAKAVDFIPVESLETTM 180
 Db 1146 GDSRGLSLSPRISYLVKSGGGPGLCPAGHAGIFRAAVCTIRGAKAVDFIPVENLETTM 1205
 QY 181 RS 182
 Db 1206 RS 1207
 RESULT 2
 POLG_HCVH STANDARD; PRT: 3011 AA.
 AC P27958;
 DT 01-AUG-1992 (Rel. 23, Created)
 DT 01-AUG-1992 (Rel. 23, Last sequence update)
 DT 15-SEP-2003 (Rel. 42, Last annotation update)
 DE Genome polyprotein [Contains: Capsid protein C (Core protein) (P22);
 DE Envelope glycoprotein E1 (GP32) (GP35); Envelope glycoprotein E2
 DE (GP68) (GP70) (NS1); protein P7; Nonstructural protein NS2 (P21)
 DE (EC 3.4.99.-); Protease/helicase NS3 (P70) (Hepacivirin)
 DE (EC 3.4.21.98); Nonstructural protein NS4A (P4); Nonstructural protein
 DE NS4B (P27); Nonstructural protein NS5A (P56); Nonstructural protein
 DE NS5B (P66) (P70) (RNA-directed RNA polymerase) (EC 2.7.7.48)].
 OS Hepatitis C virus (isolate H) (HCV).
 OC Viruses; ssRNA positive-strand viruses, no DNA stage: Flaviviridae;
 OC Hepacivirus.
 OX NCB:TaxID=11106;
 RN [1]-TaxID=11106;
 SEQUENCE FROM N.A.
 RX MEDLINE=92052256; PubMed=1658800;
 RA Inchauspé G., Zebedee S., Lee D.H., Sugitani M., Nasoff M.,
 RA Pirce A.M.;
 RI "Genomic structure of the human prototype strain H of hepatitis C
 RI virus: comparison with American and Japanese isolates.";
 RL Proc. Natl. Acad. Sci. U.S.A. 88:10292-10296(1991).
 RN [2];
 RP X-RAY CRYSTALLOGRAPHY (2.1 ANGSTROMS) OF 1207-1657.
 RX MEDLINE=97331322; PubMed=9187654;
 RA Yao N., Hesson T., Cable M., Hong Z., Kwong A.D., Le H.V., Weber P.C.;
 RL "Structure of the hepatitis C virus RNA helicase domain.";
 RN Nat. Struct. Biol. 4:463-467(1997).
 RP X-RAY CRYSTALLOGRAPHY (2.2 ANGSTROMS) OF 1192-1657.
 RX MEDLINE=9815432; PubMed=9493270;
 RA Kim J.L., Morgenstern K.A., Griffith J.P., Dwyer M.D., Thomson J.A.,
 RA Mursko M.A., Lin C., Caron P.R.;
 RI "Hepatitis C virus NS3 RNA helicase domain with a bound
 RI oligonucleotide: the crystal structure provides insights into the mode
 RI of unwinding.";
 RL Structure 6:89-100(1998).
 CC -!- FUNCTION: PROTEASE NS2 IS RESPONSIBLE FOR THE CLEAVAGE OF NS2-NS3.
 CC -!- FUNCTION: PROTEASE NS3 IS RESPONSIBLE FOR THE CLEAVAGE OF
 CC NS3-NS4A, NS4A-NS4B, NS4B-NS5A AND NS5A-NS5B.
 CC -!- FUNCTION: NS4A FORMS A COMPLEX WITH NS3 AND IS ESSENTIAL FOR THE
 CC ACTIVATION OF NS3.
 CC -!- FUNCTION: NS5A SEEMS TO HAVE A TRANSCRIPTIONAL ACTIVATORY ROLE.
 CC -!- FUNCTION: NS5B IS A RNA-DEPENDENT RNA POLYMERASE THAT PLAYS AN
 CC ESSENTIAL ROLE IN THE VIRUS REPLICATION.
 CC -!- CATALYTIC ACTIVITY: Hydrolysis of four peptide bonds in the viral
 CC precursor polyprotein, commonly with Asp or Glu in the p6
 CC position, Cys or Thr in P1 and Ser or Ala in P1'.
 CC -!- CATALYTIC ACTIVITY: N nucleoside triphosphate = N diphosphate +
 CC [RNA](N).
 CC -!- SUBUNIT: THE VIRION OF THIS VIRUS IS A NUCLEOCAPSID COVERED BY A
 CC LIPOPROTEIN ENVELOPE. THE ENVELOPE CONSISTS OF TWO PROTEINS: E1
 CC AND E2. THE NUCLEOCAPSID IS A COMPLEX OF PROTEIN C AND MRNA.


```
Matches 175: Conservative 4: Mismatches 3: Indels 0: Gaps 0:
QY 1 MAPITAYAAQTRGLGCIITSLTGKDKNOVEVOIVTSAAQTEATGTCGNCWTVYHCA 60
DB 1026 LAPITAYAAQTRGLGCIITSLTGKDKNOVEVOIVTSAAQTEATGTCGNCWTVYHCA 1085
QY 61 GRTIASPKSPVQMTYVNDKIDLVGPAHOGSRSLTPCAGSSDLYLVTRHADVPVRRR 120
DB 1086 GRTIASPKSPVQMTYVNDKIDLVGPAHOGSRSLTPCAGSSDLYLVTRHADVPVRRR 1145
QY 121 GDSRGSLLSPRISYLYKSGSGFLPCGAGHAGCZIFRAAVCTRGVAKAVDFIPVESLETIM 180
DB 1146 GDSRGSLLSPRISYLYKSGSGFLPCGAGHAGCZIFRAAVCTRGVAKAVDFIPVENLETIM 1205
QY 181 RS 182
DB 1206 RS 1207

RESULT 3
POLG_HCVTV
ID POLG_HCVTV STANDARD: PRT: 3010 AA.
AC P29846;
DT 01-APR-1993 (Rel. 25, Created)
DT 01-APR-1993 (Rel. 25, Last sequence update)
DE Genome polyprotein [contains: Capsid protein C (Core protein) (P22);
DE Envelope glycoprotein E1 (GP32) (GP35); Envelope glycoprotein E2
DE (GP68) (GP70) (NS1); Protein P7; Nonstructural protein NS2 (P21)
DE (EC 3.4.22.-); Protease/helicase NS3 (P70) (Hepacivirin)
DE (EC 3.4.21.98); Nonstructural protein NS4A (P4); Nonstructural protein
DE NS4B (P27); Nonstructural protein NS5A (P56); Nonstructural protein
DE NS5B (P66) (P70) (RNA-directed RNA polymerase) (EC 2.7.7.48)];
OS Hepatitis C virus (isolate Taiwan) (HCV).
OC Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
OC Hepacivirus.
OX NCBI_TaxID=31645;
RN [1]
RP SEQUENCE FROM N.A.
RA MEDLINE=9220206; PubMed=1314449;
RX Chen P.J., Lin M.H., Tai K.F., Liu P.C., Lin C.J., Chen D.S.:
RT "The Taiwanese hepatitis C virus genome: sequence determination and
RT mapping the 5' termini of viral genomic and antigenomic RNA.";
RL Virology 189:102-113(1993)
CC -1- FUNCTION: THE SMALL PROTEINS NS2A, NS2B, NS4A AND NS4B ARE
CC HYDROPHOBIC, SUGGESTING A POSSIBLE MEMBRANE-RELATED FUNCTION.
CC NS3 AND NS5 MAY PLAY A ROLE IN THE VIRAL RNA REPLICATION.
CC -1- CATALYTIC ACTIVITY: Hydrolysis of four peptide bonds in the viral
CC precursor polyprotein, commonly with Asp or Glu in the p6
CC position, Cys or Thr in p1 and Ser or Ala in p1'.
CC -1- CATALYTIC ACTIVITY: Nucleoside triphosphate = N diphosphate +
CC [RNAI](N).
CC -1- SUBUNIT: THE VIRION OF THIS VIRUS IS A NUCLEOCAPSID COVERED BY A
CC LIPOPROTEIN ENVELOPE. THE ENVELOPE CONSISTS OF TWO PROTEINS:
CC PROTEIN M AND GLYCOPROTEIN E. THE NUCLEOCAPSID IS A COMPLEX OF
CC PROTEIN C AND MRNA.
CC -1- SIMILARITY: THE PROTEASE BELONGS TO PEPTIDASE FAMILY S29.
CC
CC -----
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CC -----
CC EMBL: M84754; ; NOT_ANNOTATED_CDS.
CC PIR: A40244; GNMVTV.
CC PDB: 1N64; 25-FEB-03.
CC PDB: 1NS3; 08-APR-98.
CC MEROPS: S29.001; .
CC InterPro: IPR001410; DEAD.
DR InterPro: IPR002522; HCV_capsid.
DR InterPro: IPR002521; HCV_core.
DR InterPro: IPR002531; HCV_NS1.
DR InterPro: IPR002518; HCV_NS2.
DR InterPro: IPR004109; HCV_NS3.
DR InterPro: IPR000745; HCV_NS4a.
DR InterPro: IPR001490; HCV_NS4b.
DR InterPro: IPR002868; HCV_NS5a.
DR InterPro: IPR002166; HCV_RdRP.
DR InterPro: IPR007095; RNA_pol_DS_PS.
DR InterPro: IPR007094; RNA_pol_Psvir.
DR Pfam: PF01543; HCV_core; 1.
DR Pfam: PF01542; HCV_core; 1.
DR Pfam: PF01539; HCV_env; 1.
DR Pfam: PF01560; HCV_NS1; 1.
DR Pfam: PF01538; HCV_NS2; 1.
DR Pfam: PF02907; HCV_NS3; 1.
DR Pfam: PF01006; HCV_NS4a; 1.
DR Pfam: PF01001; HCV_NS4b; 1.
DR Pfam: PF01506; HCV_NS5a; 1.
DR Pfam: PF00271; Helicase_C; 1.
DR Pfam: PF00998; Viral_RdRP; 1.
DR ProDom: PD186062; HCV_NS1; 1.
DR SMARI: SM00487; DEXDC; 1.
DR PolyProtein: Glycoprotein; Transferase; RNA-directed RNA polymerase;
KW Core protein; Coat protein; Envelope protein; Helicase; ATP-binding;
KW Transmembrane; Nonstructural protein; Hydrolase; Serine protease;
KW 3D-structure.
FT INIT_MET 1 1 REMOVED FROM CAPSID PROTEIN C BY THE
FT CHAIN 1 115 CELLULAR AMINOPEPTIDASE..
FT CHAIN 116 191 CORE PROTEIN (POTENTIAL)..
FT CHAIN 192 383 MATRIX PROTEIN (POTENTIAL)..
FT CHAIN 384 729 MAJOR ENVELOPE PROTEIN E (POTENTIAL)..
FT CHAIN 730 1006 NONSTRUCTURAL PROTEIN NS1/E2 (POTENTIAL)..
FT CHAIN 1007 1615 NONSTRUCTURAL PROTEIN NS2 (POTENTIAL)..
FT CHAIN 1616 1862 PROTEASE/HELICASE NS3 (POTENTIAL)..
FT CHAIN 1863 2013 NONSTRUCTURAL PROTEIN NS4A (POTENTIAL)..
FT CHAIN 2014 3010 NONSTRUCTURAL PROTEIN NS4B (POTENTIAL)..
FT TRANSMEM 347 369 RNA-DIRECTED RNA POLYMERASE (POTENTIAL)..
FT ACT_SITE 1083 1083 POTENTIAL.
FT ACT_SITE 1107 1107 CHARGE RELAY SYSTEM (BY SIMILARITY)..
FT ACT_SITE 1165 1165 CHARGE RELAY SYSTEM (BY SIMILARITY)..
FT NP_BIND 1230 1237 ATP (POTENTIAL)..
FT SITE 1316 1319 DECH BOX.
FT CARBOHYD 196 196 N-LINKED (GLCNAC.. ) (POTENTIAL)..
FT CARBOHYD 209 209 N-LINKED (GLCNAC.. ) (POTENTIAL)..
FT CARBOHYD 233 233 N-LINKED (GLCNAC.. ) (POTENTIAL)..
FT CARBOHYD 234 234 N-LINKED (GLCNAC.. ) (POTENTIAL)..
FT CARBOHYD 250 250 N-LINKED (GLCNAC.. ) (POTENTIAL)..
FT CARBOHYD 305 305 N-LINKED (GLCNAC.. ) (POTENTIAL)..
FT CARBOHYD 417 417 N-LINKED (GLCNAC.. ) (POTENTIAL)..
FT CARBOHYD 423 423 N-LINKED (GLCNAC.. ) (POTENTIAL)..
FT CARBOHYD 430 430 N-LINKED (GLCNAC.. ) (POTENTIAL)..
FT CARBOHYD 448 448 N-LINKED (GLCNAC.. ) (POTENTIAL)..
FT CARBOHYD 532 532 N-LINKED (GLCNAC.. ) (POTENTIAL)..
FT CARBOHYD 540 540 N-LINKED (GLCNAC.. ) (POTENTIAL)..
FT CARBOHYD 556 556 N-LINKED (GLCNAC.. ) (POTENTIAL)..
FT CARBOHYD 576 576 N-LINKED (GLCNAC.. ) (POTENTIAL)..
FT CARBOHYD 623 623 N-LINKED (GLCNAC.. ) (POTENTIAL)..
FT CARBOHYD 645 645 N-LINKED (GLCNAC.. ) (POTENTIAL)..
FT CARBOHYD 2041 2041 N-LINKED (GLCNAC.. ) (POTENTIAL)..
FT CARBOHYD 2077 2077 N-LINKED (GLCNAC.. ) (POTENTIAL)..
FT CARBOHYD 2240 2240 N-LINKED (GLCNAC.. ) (POTENTIAL)..
FT CARBOHYD 2529 2529 N-LINKED (GLCNAC.. ) (POTENTIAL)..
FT CARBOHYD 2788 2788 N-LINKED (GLCNAC.. ) (POTENTIAL)..
SQ SEQUENCE 3010 AA; 327047 MW; AAD267D55CDFE215 CRC64;
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Query Match 93.6%; Score 892; DB 1; Length 3010;

Best Local Similarity 90.1%; Pred.No. 2.7e-77;

Matches 164; Conservative 12; Mismatches 6; Indels 0; Gaps 0;


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SITE 1316 1319 DECH BOX.
FT CARBOHYD 196 196 N-LINKED (GLCNAC) (POTENTIAL)
FT CARBOHYD 209 209 N-LINKED (GLCNAC) (POTENTIAL)
FT CARBOHYD 234 234 N-LINKED (GLCNAC) (POTENTIAL)
FT CARBOHYD 250 250 N-LINKED (GLCNAC) (POTENTIAL)
FT CARBOHYD 305 305 N-LINKED (GLCNAC) (POTENTIAL)
FT CARBOHYD 417 417 N-LINKED (GLCNAC) (POTENTIAL)
FT CARBOHYD 423 423 N-LINKED (GLCNAC) (POTENTIAL)
FT CARBOHYD 430 430 N-LINKED (GLCNAC) (POTENTIAL)
FT CARBOHYD 448 448 N-LINKED (GLCNAC) (POTENTIAL)
FT CARBOHYD 532 532 N-LINKED (GLCNAC) (POTENTIAL)
FT CARBOHYD 540 540 N-LINKED (GLCNAC) (POTENTIAL)
FT CARBOHYD 556 556 N-LINKED (GLCNAC) (POTENTIAL)
FT CARBOHYD 576 576 N-LINKED (GLCNAC) (POTENTIAL)
FT CARBOHYD 623 623 N-LINKED (GLCNAC) (POTENTIAL)
FT CARBOHYD 645 645 N-LINKED (GLCNAC) (POTENTIAL)
FT CARBOHYD 2041 2041 N-LINKED (GLCNAC) (POTENTIAL)
FT CARBOHYD 2077 2077 N-LINKED (GLCNAC) (POTENTIAL)
FT CARBOHYD 2240 2240 N-LINKED (GLCNAC) (POTENTIAL)
FT CARBOHYD 2529 2529 N-LINKED (GLCNAC) (POTENTIAL)
FT CARBOHYD 2788 2788 N-LINKED (GLCNAC) (POTENTIAL)
FT STRAND 1031 1035 N-LINKED (GLCNAC) (POTENTIAL)
FT HELIX 1039 1047
FT STRAND 1050 1050
FT STRAND 1059 1063
FT STRAND 1068 1074
FT TURN 1075 1076
FT STRAND 1077 1081
FT HELIX 1082 1085
FT TURN 1086 1087
FT STRAND 1090 1092
FT TURN 1093 1094
FT STRAND 1095 1097
FT STRAND 1101 1103
FT TURN 1104 1107
FT STRAND 1108 1112
FT STRAND 1120 1120
FT STRAND 1122 1122
FT STRAND 1129 1133
FT TURN 1135 1136
FT STRAND 1139 1144
FT STRAND 1149 1157
FT HELIX 1158 1161
FT TURN 1162 1163
FT TURN 1165 1166
FT STRAND 1168 1171
FT TURN 1172 1174
FT STRAND 1175 1186
FT TURN 1187 1188
FT STRAND 1189 1197
FT HELIX 1198 1202
FT TURN 1203 1204
FT STRAND 1205 1208
SQ SEQUENCE 3010 AA: 327165 MW: F942255ECDFD9C3RC64;

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Query March 93.5% Score 991: 58 11 length 3010;
 Rest Local Similarity 89.0% Pred.No. 3.4e-77;
 Matches 162; Conservative 15; Mismatches 5; Indels 0; Gaps 0;

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Qy 1 MAPITAYAQTRG:LGCIITSLTRCKNQVEGEVGVISIAACTFLATCINGVCWTVYHGA 50
Db 1026 LAPITAYSQTRG:LGCIITSLTRCKNQVEGEVGVQVWSTATQSFAPATCVNGVCWTVYHGA 1085
Qy 61 GTRITASPKGFVIMYTNVDRKLVGVGAPQGSRSLSPTCTGSGSDLYLVTRHADVPVRRR 120
Db 1086 GSKTLAPKPGPITOMY:NVQDLVGVWPKPGCARSLTFCICGSSDLYLVTRHADVPVRRR 1145
Qy 12: GDSRGSLLSPRPISY:JAGSGGGPLICVAGHAGVFRAAVCTRGVAKAVDFIPVESLETTM 180
Db 1146 GDSRGSLLSPRPVSYERKSGGGPLLCFPGHAGVIFRAAVCTRGVAKAVDFIPVESNETTM 1205
Qy 181 RS 182

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UB 1206 RS 1207
RESULT 5
POLG_HCV7A
ID POLG_HCV7A STANDARD: PRT: 3010 AA.
AC P26662;
DT 01-AUG-1992 (Rel. 23, Created)
DT 01-AUG-1992 (Rel. 23, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Genome polyprotein [Contains: Capsid protein C (Core protein) (P22);
DE Envelope glycoprotein E1 (GP32) (GP35); Envelope glycoprotein E2
DE (GP68) (GP70) (NS1); Protein P7; Nonstructural protein NS2 (P21)
DE (EC 3.4.22.-); Protease/helicase NS3 (P70) (Hepacivirin)
DE (EC 3.4.21.98); Nonstructural protein NS4A (P4); Nonstructural protein
DE NS4B (P27); Nonstructural protein NS5A (P56); Nonstructural protein
DE NS5B (P66) (P70) (RNA-directed RNA polymerase) (HCV).
OS Hepatitis C virus (isolate Japanese) (HCV).
OC Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
OC Hepacivirus.
OX NCBI_TaxID=11116;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=9108550; PubMed=2175903;
RA Kato N., Hijikata M., Ootsuyama Y., Nakagawa M., Ohkoshi S.,
RA Sugimura T., Shimotohno K.;
RT "Molecular cloning of the human hepatitis C virus genome from
RT Japanese patients with non-A, non-B hepatitis.";
RL Proc. Natl. Acad. Sci. U.S.A. 87:9524-9528(1990).
RN [2]
RP DISCUSSION OF SEQUENCE.
RX MEDLINE=91192160; PubMed=1849488;
RA Kato N., Hijikata M., Nakagawa M., Ootsuyama Y., Muraiso K.,
RA Ohkoshi S., Shimotohno K.;
RT "Molecular structure of the Japanese hepatitis C viral genome.";
RL FEBS Lett. 280:325-328(1991).
CC -!- FUNCTION: THE SMALL PROTEINS NS2A, NS2B, NS4A AND NS4B ARE
CC HYDROPHOBIC, SUGGESTING A POSSIBLE MEMBRANE-RELATED FUNCTION.
CC NS3 AND NS5 MAY PLAY A ROLE IN THE VIRAL RNA REPLICATION.
CC -!- CATALYTIC ACTIVITY: Hydrolysis of four peptide bonds in the viral
CC precursor polyprotein, commonly with Asp or Glu in the P6
CC position, Cys or Thr in P1 and Ser or Ala in P1'.
CC -!- CATALYTIC ACTIVITY: N nucleoside triphosphate - N diphosphate +
CC {RNA}(N).
CC -!- SUBUNIT: THE VIRION OF THIS VIRUS IS A NUCLEOCAPSID COVERED BY A
CC LIPOPROTEIN ENVELOPE. THE ENVELOPE CONSISTS OF TWO PROTEINS:
CC PROTEIN M AND GLYCOPROTEIN E. THE NUCLEOCAPSID IS A COMPLEX OF
CC PROTEIN C AND RNA.
CC -!- SIMILARITY: THE PROTEASE BELONGS TO PEPTIDASE FAMILY S29.
CC
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CC
CC EMBL: D90208; BAAL4233.1; -
CC PIR: A35253; GNWVCL.
CC HSP: P26663; LUXP.
CC MEROPS: S29.001; -
CC MEROPS: U39.001; -
CC
CC InterPro: IPR001410; DEAD.
CC InterPro: IPR002522; HCV_capsid.
CC InterPro: IPR002521; HCV_core.
CC InterPro: IPR002519; HCV_env.
CC InterPro: IPR002531; HCV_NSI.
CC InterPro: IPR002518; HCV_NS2.
CC InterPro: IPR004109; HCV_NS3.
CC InterPro: IPR000745; HCV_NS4a.
CC InterPro: IPR001490; HCV_NS4b.
CC InterPro: IPR002868; HCV_NS5a.
CC

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DR InterPro: IPR002166; HCV RdRp.
 DR InterPro: IPR001650; Helicase_C.
 DR InterPro: IPR007095; RNA_pol_DS_PS.
 DR InterPro: IPR007094; RNA_pol_PSVir.
 DR Pfam: PF01543; HCV_capsid; 1.
 DR Pfam: PF01542; HCV_core; 1.
 DR Pfam: PF01539; HCV_env; 1.
 DR Pfam: PF01560; HCV_NS1; 1.
 DR Pfam: PF01538; HCV_NS2; 1.
 DR Pfam: PF02907; HCV_NS3; 1.
 DR Pfam: PF01006; HCV_NS4a; 1.
 DR Pfam: PF01001; HCV_NS4b; 1.
 DR Pfam: PF01506; HCV_NS5a; 1.
 DR Pfam: PF00271; Helicase_C; 1.
 DR Pfam: PF00998; Viral_RdRp; 1.
 DR ProDom: PD186662; HCV_NS1; 1.
 DR SMART: SM00487; DDXC; 1.
 KW Polyprotein: Glycoprotein; Transferrase; RNA-directed RNA polymerase;
 KW Core protein; Envelope protein; Helicase; ATP-binding;
 KW Transmembrane; Nonstructural protein; Hydrolase; Serine protease.
 FT INIT_MET 1
 FT CHAIN 1 125
 FT CHAIN 1-5 194
 FT CHAIN 192 383
 FT CHAIN 384 729
 FT CHAIN 730 1026
 FT CHAIN 1027 1615
 FT CHAIN 1616 1862
 FT CHAIN 1863 2013
 FT CHAIN 2014 3012
 FT CHAIN 3013 369
 FT TRANSMEM 1
 FT ACT_SITE 1083 1083
 FT ACT_SITE 1107 1107
 FT ACT_SITE 1155 1155
 FT NP_BIND 1237 1237
 FT SITE 1316 1319
 FT CARBOHYD 135 136
 FT CARBOHYD 209 209
 FT CARBOHYD 234 234
 FT CARBOHYD 250 250
 FT CARBOHYD 305 305
 FT CARBOHYD 417 417
 FT CARBOHYD 423 423
 FT CARBOHYD 430 430
 FT CARBOHYD 448 448
 FT CARBOHYD 532 532
 FT CARBOHYD 556 556
 FT CARBOHYD 576 576
 FT CARBOHYD 623 623
 FT CARBOHYD 645 645
 FT CARBOHYD 2041 2041
 FT CARBOHYD 2077 2077
 FT CARBOHYD 2240 2240
 FT CARBOHYD 2788 2788
 FT SEQUENCE 3010 AA: 327017 MW: AA993794F463B185 CMC84;
 Query Match 93.1%; Score 887; DB 1; Length 3010;
 Best Local Similarity 87.9%; Pred. No. 8, 2e-77;
 Matches 160; Conservative 17; Mismatches 5; Indels 0; Gaps 0;
 1 MAPITAYAOQTGRLCCITTSITGRKKNQVEVQVSTAACTFLATGNGCVTWYHCA 60
 :|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
 1026 LAPITAYSOOTRGLGCHITTSITGRKKNQVDQVQLSTATQSFATCVNGVGVYHGA 1085
 :|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
 61 GTRTASPKGPVQWYTNVDKLVGKAPQGSSESLTCTCGSSDLYLVTRHADVVRKR 120
 :|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
 1086 GSKTLAGKGPQIQWYTNVDQLVGVWAPPGRASHPICCGSSDLYLVTRHADVVRKR 1145
 :|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
 121 GDSRGLSPRPISYIKGSSGCPGLLCPAGHAYGIFRAAVCTRGVAKVDFIPVESMETM 180
 :|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
 1146 GDSRGLSPRPISYIKGSSGCPGLLCPGSHVVGIFRAAVCTRGVAKVDFIPVESMETM 1205

QY 181 RS 182
 II
 DB 1206 RS 1207
 RESULT 6
 POLQ_HCVJT STANDARD: PRT: 30-0 AA.
 ID POLQ_HCVJT
 AC Q00269;
 DT 01-APR-1993 (Rel.: 25, Created)
 DT 01-APR-1993 (Rel.: 25, Last sequence update)
 DT 15-SEP-2003 (Rel.: 42, Last annotation update)
 SE Genome polyprotein [Contains: Capsid protein C (Core protein) (P22);
 DE Envelope glycoprotein E1 (GP32) (GP35); Envelope glycoprotein E2
 DE (GP68) (GP70) (NS1); Protein P7; Nonstructural protein NS2 (P21)
 DE (EC 3.4.22.-); Protease/helicase NS3 (P70) (Hepacivirin)
 DE (EC 3.4.21.98); Nonstructural protein NS4A (P4); Nonstructural protein
 DE NS4B (P27); Nonstructural protein NS5A (P56); Nonstructural protein
 DE NS5B (P66) (P70) (RNA-directed RNA polymerase) (EC 2.7.7.48)].
 OS Hepatitis C virus (isolate HC-JT) (HCV).
 OC Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
 OC Hepacivirus.
 OC NCBI_TaxID=31642;
 RN SEQUENCE FROM N.A.
 RN [1]
 RX MEDLINE=92295714; PubMed=1318627;
 RA Tanaka T., Kato N., Nakagawa M., Gotsuyama Y., Cho M.C.,
 RA Nakazawa T., Hjikata M., Ishimura Y., Shimotohno K.;
 RT "Molecular cloning of hepatitis C virus genome from a single Japanese
 RT carrier: sequence variation within the same individual and among
 RT infected individuals.";
 RL Virus Res. 23:39-53(1992).
 CC -!- FUNCTION: THE SMALL PROTEINS NS2A, NS2B, NS4A AND NS4B ARE
 CC HYDROPHOBIC, SUGGESTING A POSSIBLE MEMBRANE-RELATED FUNCTION.
 CC NS3 AND NS5 MAY PLAY A ROLE IN THE VIRAL RNA REPLICATION.
 CC -!- CATALYTIC ACTIVITY: Hydrolysis of four peptide bonds in the viral
 CC precursor polyprotein, commonly with Asp or Glu in the P6
 CC position, Cys or Thr in P1 and Ser or Ala in P1'.
 CC -!- CATALYTIC ACTIVITY: N nucleoside triphosphate -> N diphosphate +
 CC RNA(N).
 CC -!- SUBUNIT: THE VIRION OF THIS VIRUS IS A NUCLEOCAPSID COVERED BY A
 CC LIPOPROTEIN ENVELOPE. THE ENVELOPE CONSISTS OF TWO PROTEINS:
 CC PROTEIN M AND GLYCOPROTEIN E. THE NUCLEOCAPSID IS A COMPLEX OF
 CC PROTEIN C AND MRNA.
 CC -!- SIMILARITY: THE PROTEASE BELONGS TO PEPTIDASE FAMILY S29.
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 DR EMBL: D11168; BA01943.1;
 DR PIR: A45573; A45573.
 DR PDB: 1A1Q; 25-MAR-98.
 DR PDB: 1JXP; 14-JAN-98.
 DR MFROPS: S29.001;
 DR MFROPS: U39.001;
 DR InterPro: IPR001410; DEAD.
 DR InterPro: IPR002522; HCV_capsid.
 DR InterPro: IPR002521; HCV_core.
 DR InterPro: IPR002519; HCV_env.
 DR InterPro: IPR002531; HCV_NS1.
 DR InterPro: IPR002518; HCV_NS2.
 DR InterPro: IPR004109; HCV_NS3.
 DR InterPro: IPR000745; HCV_NS4a.
 DR InterPro: IPR002868; HCV_NS4b.
 DR InterPro: IPR002166; HCV_RdRp.
 DR InterPro: IPR007095; RNA_pol_DS_PS.
 DR InterPro: IPR007094; RNA_pol_PSVir.

DR Pfam: PF02907; HCV_NS3; 1.
 DR Pfam: PF01006; HCV_NS4a; 1.
 DR Pfam: PF01901; HCV_NS4b; 1.
 DR Pfam: PF01536; HCV_NS5a; 1.
 DR Pfam: PF00271; helicase_C; 1.
 DR Pfam: PF00598; Viral_RDRP; 1.
 DR ProDom: PD185062; HCV_NS1; 1.
 DR SMART: SM00487; DEXDC; 1.
 KW Polypeptide; Glycoprotein; Transferase; RNA-directed RNA polymerase;
 KW Core protein; Coat protein; Envelope protein; Helicase; ATP-binding;
 KW Transmembrane; Nonstructural protein; Hydrolase; Serine protease;
 FT INIT_MET 1
 FT CHAIN 1 125
 FT CHAIN 116 191
 FT CHAIN 192 393
 FT CHAIN 384 733
 FT CHAIN 734 1010
 FT CHAIN 1011 1619
 FT CHAIN 1620 1856
 FT CHAIN 1857 2017
 FT CHAIN 2018 3033
 FT TRANSMEM 347 369
 FT ACT_SITE 1087 1087
 FT ACT_SITE 1111 1111
 FT ACT_SITE 1169 1169
 FT NP_BIND 1234 1241
 FT SITE 1320 1323
 FT CARBOHYD 196 196
 FT CARBOHYD 209 209
 FT CARBOHYD 234 234
 FT CARBOHYD 305 305
 FT CARBOHYD 417 417
 FT CARBOHYD 423 423
 FT CARBOHYD 436 436
 FT CARBOHYD 448 448
 FT CARBOHYD 477 477
 FT CARBOHYD 534 534
 FT CARBOHYD 542 542
 FT CARBOHYD 558 558
 FT CARBOHYD 578 578
 FT CARBOHYD 627 627
 FT CARBOHYD 649 649
 FT CARBOHYD 1091 1091
 FT CARBOHYD 2038 2038
 FT CARBOHYD 2811 2811
 SQ SEQUENCE 3033 AA; 329155 MW; F9575C1A273BE9E CAC64;
 Query Match 74.7%; Score 712; DR 1; Length 3033;
 Best Local Similarity 69.8%; Pred. No. 53e-60;
 Matches 127; Conservative 29; Mismatches 26; Indels 0; Gaps 0;
 QY 1 MAPITAYAOQTGGLGCTTSLTGKDNQVEGFVOIVSTAAQTFLATPCINGVGVWVYHCA 60
 Db 1030 LAPITAYAOQTGGLGCTTSLTGKDNQVEGFVOIVSTAAQTFLATPCINGVGVWVYHCA 1069
 QY 61 GTRTIASPKGPVIMYNTVNDKLVGKAPQGSRLTPCTCGSSDLXLYVTHAVIPVRRR 120
 Db 1090 GNKTLAGSRGPVTOMYSAAEGDLVGMWSPPTKSLPCTCGSSDLXLYVTHAVIPVRRR 1149
 QY 121 GDSRGSLLSPRISLYLSSGSGGPGLLCPAGHAGVIFRAAVCTRGVAKAVDFIPVESLET 180
 Db 1150 GDRKCALLSPRLSTLKGSGGPGVLCPRHAGVIFRAAVCTRGVAKAVDFIPVESLET 1209
 QY 181 RS 182
 Db 1210 RS 1211
 RESULT 9
 ID PAAD_PSEAE STANDARD; PRI: 209 AA.
 AC C9HX08;

DT 16-OCT-2001 (Rel. 40, Created)
 DT 16-OCT-2001 (Rel. 40, Last sequence update)
 DT 16-OCT-2001 (Rel. 40, Last annotation update)
 DE Probable aromatic acid decarboxylase (EC 4.1.1.1).
 GN PA4019.
 OS Pseudomonas aeruginosa.
 CC Bacteria; Proteobacteria; Gammaproteobacteria; Pseudomonadales;
 CC Pseudomonadaceae; Pseudomonas.
 CX NCBI_TaxID=287;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=ATCC 15692 / PAO.;
 RX MEDLINE=20437337; PubMed=10984043;
 RA Stover C.K., Pham X.-Q.T., Erwin A.L., Mizoguchi S.D., Warren P.,
 RA Hickey M.J., Brinkman F.S.L., Hufnagle D.O., Kowalik D.J., Lagrou M.,
 RA Garber R.J., Goltry L., Tolentino E., Westbrock-Wadman S., Yuan Y.,
 RA Brody L.L., Coulter S.N., Folger K.R., Kas A., Larbig K., Lim R.M.,
 RA Smith K.A., Spencer D.H., Wong G.K.-S., Wu Z., Paulsen I.T.,
 RA Reizer J., Saiter M.H., Hancock R.E.W., Lory S., Olson M.V.;
 RI "Complete genome sequence of Pseudomonas aeruginosa PAO1, an
 RI opportunistic pathogen";
 RL Nature 406:959-964 (2000).
 CC -1- SIMILARITY: BELONGS TO THE POLYPRENYL P-HYDROXYBENZOATE /
 CC PHENYLACRYLIC ACID DECARBOXYLASES FAMILY.
 CC -----
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 CC or send an email to license@sib-sib.ch).
 CC -----
 CC EMBL: AF004618; AAG07406.1; .
 DR PIR: H83144; H83144.
 DR InterPro: IPR003382; Flavoprotein.
 DR Pfam: PF02441; Flavoprotein: 1.
 KW Hypothetical protein; lyase; Decarboxylase; Complete proteome.
 SQ SEQUENCE 209 AA; 22367 MW; G1FD081CC495D3F6 CRC64;
 Query Match 9.1%; Score 87; DR 1; Length 209;
 Best Local Similarity 26.2%; Pred. No. 0.28;
 Matches 55; Conservative 18; Mismatches 61; Indels 76; Gaps 12;
 QY 8 AQQTGGLGCTTSLTGKDNQVEGFVOIVSTAAQTFLATPCINGVGVWVYHCA 66
 Db 17 AQVGLRLDCLV-----QFEVHFPLISKAAQLVMAI-----ETVVA 53
 QY 67 SPKGP-----VQMVTNVDKLVGKAPQGSRLTP-----CICGSSDL 105
 Db 54 LPKAPQAMQAFLETGGAAGQIRVFGND-----WMAPPASGSGAPNAWICPSTGTL 108
 QY 106 -----YLVRGADVIPVRRGDSRGSLLSPR--PTS-----YKGGSGGPGLLCPA 148
 Db 109 SAVATGACNKLIERADVALKER---RPLVLVPEAFPSHLENMILKSLNLGAVILPA 164
 QY 149 GHAVGIFRAAVCTRGVAKAVDFIPVESLET 178
 Db 165 --APGFYHQ---PQSVEDLVDFVVAIINT 189
 RESULT 10
 ID HHOA_ARATH STANDARD; PRI: 321 AA.
 AC Q9SEL7; O49507;
 DT 16-OCT-2001 (Rel. 40, Created)
 DT 16-OCT-2001 (Rel. 40, Last sequence update)
 DT 15-SEP-2003 (Rel. 42, Last annotation update)
 DE protease HhoA, chloroplast precursor (EC 3.4.21.-).
 GN HHOA OR AT4G18370 OR F28J12.30.
 OS Arabidopsis thaliana (Mouse-ear cress).
 CC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 CC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicotyledons; Rosidae;

CC extroids II; Brassicales; Brassicaceae; Arabidopsis.
 CC NCBI_TaxID=3702;
 RN [1]
 RP SEQUENCE FROM N.A.
 RA Lersch M.H.A., Sokolenko A., Herrmann R.G.:
 RI "Identification and characterization of the chloroplast HsA protease,
 RI a homolog to the bacterial periplasmic protease HsA.",
 RI Submitted (DEC-1998) to the EMBL/GenBank/DBJ databases.
 RN [2]
 RP SEQUENCE FROM N.A.
 RC STRAIN=cv. Columbia;
 RX MEDLINE=26083488; PubMed=10617198;
 RA Mayer K.F.A., Schueller C., Wambutt R., Murphy G., Volckaert G.,
 RA Pohl T., Duesterhoeft A., Stiekema W., Ettian K.-D., Terryn M.,
 RA Harris B., Ansoore W., Brandt P., Grivell L., Kieger M.,
 RA Weichselgartner M., de Simone V., Obermaier B., Wache K., Mueller M.,
 RA Kreis M., Delsen M., Pulgomech P., Watson M., Schmidkeini I.,
 RA Reichert B., Portetelle E., Perez-Alonso M., Souty M., Bancroft T.,
 RA Vos P., Nonheisel C., Zimmermann W., Wadler H., Fildes P.,
 RA Langham S.-A., McCullagh B., Bilham L., Robben J.,
 RA Van der Schueren J., Grymprocz B., Chuang Y.-J., Vandenbussche P.,
 RA Bracken M., Weltyons J., Voet M., Bastiaens J., Aert R., DeCoor E.,
 RA Weitzenecker T., Bothe G., Ransperger J., Hilbert H., Braun M.,
 RA Holzer E., Brandt A., Peters S., van Staveren M., Dirkse W.,
 RA Moolman P., Klein Lankhorst R., Rose M., Hauf J., Koetter P.,
 RA Berner S., Hempel S., Fedpausch M., Lambert S., Van den Daele H.,
 RA De Keyser A., Huysschaert C., Gielon J., Villarroel R., De Clercq R.,
 RA Van Montagu M., Kogers C., Cronin A., Qaill M., Bray-Allen S.,
 RA Clark S., Dorgett J., Hall S., Kay M., Leonard N., McLay K., Mayes R.,
 RA Pettelt A., Rajandream M.A., Lyne M., Boncs V., Rechmann S.,
 RA Borkova D., Bloeker A., Scharie M., Gram M., Moehner T.-H.,
 RA Dose S., de Haan M., Maarse A., Schaefer M., Mueller-Auer S.,
 RA Gabel C., Fuchs M., Fartmann B., Grandrath K., Dauner D., Herzi A.,
 RA Neumann S., Argirou A., Vitale D., Liguori R., Piravandi E.,
 RA Massenot O., Quigley F., Clabaud G., Kuendlein A., Felber R.,
 RA Schnabl F., Hiller R., Schmidt W., Lecharny A., Abouq S.,
 RA Chedrol S., Cooke R., Berger C., Monfort A., Casacuberta E.,
 RA Gibbons T., Weber N., Vandenbol M., Barques M., Terol C., Torres A.,
 RA Perez-Perez A., Purnelle B., Best E., Johnson S., Tabor D., Jesse T.,
 RA Heijnen L., Schwarz S., Scholler P., Heber S., Francis P., Bieleke C.,
 RA Frishman D., Haase D., Lemcke K., Meves H.-W., Stocker S.,
 RA Zaccaria P., Bevan M., Wilson R.K., de la Bastide M., Habermann K.,
 RA Parnell L., Dedhia N., Grol L., Schutz K., Huang E., Spiegel L.,
 RA Sehkon M., Murray J., Sheet P., Cordes M., Abu-Thelidh J.,
 RA Stoneking T., Kalicki J., Graves T., Harmon G., Edwards J.,
 RA Latreille P., Courtney L., Cloud J., Abbott A., Scott K., Johnson D.,
 RA Minx P., Bentley D., Fulton B., Millier N., Greco T., Kemp K.,
 RA Kramer J., Fulton L., Mardis E., Dante M., Pepin K., Hillier L.,
 RA Nelson J., Spieth J., Ryan E., Andrews S., Geisel C., Layman D.,
 RA Du H., Ali J., Berghoff A., Jones K., Drone K., Cotton M., Joshi C.,
 RA Antoniou B., Zidanic M., Strong C., Sun H., Iamar H., Yordan C.,
 RA Ma P., Zhong J., Preston R., Vil D., Shekter M., Marero A., Shah A.,
 RA Swaby I.K., O'Shaughnessy A., Rodriguez M., Hoffman J., Tili S.,
 RA Granat S., Shohdy N., Hasegawa A., Hameed A., Lodhi M., Johnson A.,
 RA Chen E., Maria M., Martensen R., McCoy W.R.:
 RI "Sequence and analysis of chromosome 4 of the plant Arabidopsis
 RI thaliana."
 RL Thaliana.
 RN [3]
 RP SEQUENCE OF 72-82; 96-110; 150-159; 178-211 AND 306-320.
 RA Schubert M., Peterson U., Funk C., Haas B., Schroeder W.P.,
 RA Kieselbach T.:
 RI "The chloroplast lumen from Arabidopsis thaliana."
 RL Submitted (JUL-2001) to the SWISS-PROT data bank.
 CC -1- SUBCELLULAR LOCATION: Chloroplast; within the thylakoid lumen.
 CC -1- SIMILARITY: BELONGS TO PEPTIDASE FAMILY S2C.
 CC -1- CAUTION: Ref.2 sequences differ from that shown due to erroneous
 CC gene model prediction. AT4G18370 and AT4G18375 were originally
 CC fused into a single gene.
 CC -----
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 CC -----
 CC EMBL: AF114386; AAF24060.1; -;
 CC EMBL: AL021710; CAA16717.1; ALT_SEQ.
 CC EMBL: AL161548; CAB78839.1; ALT_SEQ.
 CC MROPS: S01.279; -;
 CC InterPro: IPR001940; ProteaseG2C.
 CC InterPro: IPR001254; Ser_protease_Try.
 CC Pfam: PF00089; trypsin; 1;
 CC PRINTS: PR00834; PROTEASES2C.
 CC Hydrolyase; Serine protease; Chloroplast; Thylakoid; Transit peptide.
 KW CHLOROPLAST (POTENTIAL).
 FT TRANSIT 1 26
 FT THYLAKOID. 71
 FT CHAIN 72 321
 FT POLY-GLU 77 87
 FT DOMEIN 77 87
 FT ACT_SITE 145 145
 FT ACT_SITE 185 186
 FT ACT_SITE 264 264
 FT ACT_SITE 40 40
 FT CONFLICT R -> G (IN REF. 1).
 SQ SEQUENCE 321 AA; 34691 MW; 66DB81E0BD27A7A7 CRC64;
 Query Match 8.8%; Score 84; DB 1; Length 321;
 Best Local Similarity 22.28; Pred. No. 0.88;
 Matches 48; Conservative 26; Mismatches 60; Indels 82; Gaps 11;
 QY 22 LTGRDKNQVEGVOLVSTAAQIFLATCINGVCM-----TVYH----- 58
 DB 117 LTDEENKIKETG-----SGVWOKLGHIVTNVHVIAKLATQFGLORCK 161
 QY 59 -----GAGTRTASPKGVHVIOMYTNVDKLVGPAPQGSRLTCTCGSSDLYLVTRHAD 113
 DB 162 VSLVDKAGTR--FSKSKIVCL--DPNDLAVLKIEITGRELNPVLTSDNLVWGSCF 217
 QY 114 VIPVRGRDSRG-----SLSPRISYUK-----GSSGGPLLCPA 148
 DB 218 AI-----GNPYGYENTLTIGVSGIGREIPSPKISSEAIQTADINSNGSGPLLDJ 272
 QY 149 GHVNGIFRAAVCIK--GVAKAVDF-IPVESLETIM 180
 DB 273 GHTIGV-NTATFTRRGSGVGNVPAIPDTVVRIV 307
 RESULT 1;
 AAMP_HUMAN
 ID AAMP_HUMAN STANDARD; PRT; 452 AA.
 AC Q13685;
 DT 15-JUL-1998 (Rel. 36, Created)
 DT 15-JUL-1998 (Rel. 36, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE Angio-associated migratory cell protein.
 GN AAMP.
 OS Homo sapiens (Human).
 CC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 CC Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
 CX NCBI_TaxID=9606;
 RN [1]
 RP SEQUENCE FROM N.A., AND PARTIAL SEQUENCE.
 RC ISSUE=Brain;
 RX MEDLINE=95262124; PubMed=7743515;
 RA Beckner M.E., Krutzsch H.C., Stracke M.L., Williams S.T.,
 RA Gallardo J.A., Liotta L.A.;
 RI Identification of a new immunoglobulin superfamily protein expressed
 RI in blood vessels with a heparin-binding consensus sequence.*;
 RL Cancer Res. 55:2140-2149(1995).
 CC -1- FUNCTION: MAY HAVE A FUNCTION IN MIGRATING CELLS.
 CC -1- TISSUE SPECIFICITY: EXPRESSED IN BLOOD VESSELS. STRONGLY EXPRESSED
 CC IN ENDOTHELIAL CELLS, CYTOTROPHOBLASTS, AND POORLY DIFFERENTIATED
 CC COLON ADENOCARCINOMA CELLS FOUND IN LYMPHATICS.
 CC -1- SIMILARITY: Contains 8 WD repeats.
 CC -----

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CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (See <http://www.isb-sib.ch/announcement/>
CC or send an email to license@isb-sib.ch).
CC -----
CC EMBL: M95627; AAA69889.1; --
CC Genew: HGNC:18; AAMP.
CC MM: 603488.
CC GO: GO:0008201; Fibrinogen binding activity; TAS.
CC InterPro: IPR001680; WD40.
CC Pfam: PF00400; WD40; 8.
CC SMART: SM00320; WD40; 8.
CC PROSITE: PS00678; WD_REPEATS_1; 1.
CC PROSITE: PS50082; WD_REPEATS_2; 6.
CC PROSITE: PS50294; WD_REPEATS_REGION; 1.
CC Repeat: WD repeat.
CC -----
FT DOMAIN 14 18 HEPARIN-BINDING (POTENTIAL).
FT DOMAIN 71 77 POLY-GLU.
FT REPEAT 107 138
FT REPEAT 150 180 WD 1.
FT REPEAT 190 220 WD 2.
FT REPEAT 231 261 WD 3.
FT REPEAT 275 306 WD 4.
FT REPEAT 333 363 WD 5.
FT REPEAT 374 404 WD 6.
FT REPEAT 416 446 WD 7.
FT REPEAT 452 485 WD 8.
SQ SEQUENCE 452 AA; DA1413D25E2B236C0 CRC64;

Query Match 8.6%; Score 82; DB 1; Length 452;

Best Local Similarity 25.3%; Pred. No. 2;

Matches 42; Conservative 13; Mismatches 47; Indels 64; Caps 9;

QY 54 WTYYHAGRTIASPKGPVQKVTNVDKLVGPAQGSRL-----TPCTCGSSDLYLV 108
DB 197 WNEWH-----PRPVULAGT-ADGNWMMKVPNGCKTFQGNCPAICGR----- 240
QY 109 TRHADVIVPRR-----GSRGSS-----LLSPRTSYLKGSSQ--GPLCPA----- 148
DB 241 -----VLPDGKRAVGVYEDGCTRIWDL-KQSGP-HVLKGTGHGPGPLTCVAANGDGLILT 295
QY 149 -----GHAVGIFR-----AAVCTRVAKAVDFPVESL 176
DB 296 GSVDCQAKLVSAI:GKVVGVFRPETVASQPSLSEGESESNVSLS 341

RESULT 12

ID Y136_TREPA STANDARD; PRT: 425 AA.
AC OB1172;
DT 16-OCT-2001 (Rel. 40, Created)
DI 16-OCT-2001 (Rel. 40, Last sequence update)
DI 16-OCT-2001 (Rel. 40, Last annotation update)
DE Hypothetical lipoprotein TP0136 precursor.
GN TP0136.
OS Treponema pallidum.
OC Bacteria; Spirochaetes; Spirochaetales; Spirochaetaceae; Treponema.
OX NCBI_Taxid=160;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=Nichols;
RX MEDLINE=98332770; PubMed=9655876;
RA Fraser C.M., Norris S.J., Weinstock G.M., White C., Sutton G.G.,
RA Dodson R., Gwin M., Hickey E.K., Clayton R., Ketchum K.A.,
RA Sodergren E., Hardham J.M., McLeod M.P., Salzberg S., Peterson J.,
RA Khalak H., Richardson D., Howell J.K., Chidambaram M., Utterback T.,
RA McDonald L., Artlich P., Bowman C., Cotton M.D., Fujii C., Garland S.,
RA Hatch B., Horst K., Roberts K., Sandusky M., Weidman J., Smith H.O.,
RA Venter J.C.;
*Complete genome sequence of Treponema pallidum, the syphilis

RT spirochete.";
RI Science 281:375-388(1998).
CC -!- SURCELLULAR LOCATION: Attached to the membrane by a lipid anchor
CC (Potential).
CC -!- SIMILARITY: BELONGS TO THE TP013X FAMILY OF LIPOPROTEINS.
CC -----
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CC or send an email to license@isb-sib.ch).
CC -----
CC EMBL: AE001199; AAC65137.1; ALT_INIT.
CC TIGR: TP0136; --
CC Hypothetical protein; Lipoprotein; Membrane; Signal;
CC Complete proteome.
FT SIGNAL 1 23 POTENTIAL.
FT CHAIN 24 485 HYPOTHETICAL LIPOPROTEIN TP0136.
FT LIPID 24 24 N-ACYL DIGLYCERIDE (POTENTIAL).
FT DOMAIN 164 178 GLY/SER-RICH.
FT DOMAIN 196 210 GLY/SER-RICH.
FT DOMAIN 253 267 GLY/SER-RICH.
FT DOMAIN 318 327 POLY-SER.
FT DOMAIN 444 447 POLY-SER.
SQ SEQUENCE 485 AA; 48984 MW; C7A4CEDC7DC5CED CRC64;
Query Match 8.6%; Score 82; DB 1; Length 485;
Best Local Similarity 24.2%; Pred. No. 2,2;
Matches 44; Conservative 13; Mismatches 65; Indels 60; Gaps 8;
QY 23 TGRDKNVEGEVQIVSTAAQTFLATCI--NGVWTVYHGAG---TRTIASPKGPVQKVT 77
DB 86 TSDSK-----KVMSTADIGNTFVLACVPGVIGYKHCNVAGAGSSSTGTGTASPSTETCSQA 140
QY 78 NVYDKLVG-----WPAGQSRSLPTCTC-----GSSDLVLTNRADVIP-----VR 118
DB 141 T-----LVGTSKPKVLVPGGTGNNCGGGGGGGSSSSSSSSCHILWLVPGGTGNNKCG 196
QY 119 RRGDSRGSLSPRLSYLK-----CGGGGGGGGGGGGGGGGGGGGGGGGGGGGGGGGGGGGG 151
DB 147 CGGGGGGGSS 256
QY 152 VG 153
DB 257 GG 258
RESULT 13
DEG_ARATH
ID DEG_ARATH STANDARD; PRT: 437 AA.
AC Q22609; O91K85;
DI 16-OCT-2001 (Rel. 40, Created)
DI 16-OCT-2001 (Rel. 40, Last sequence update)
DI 28-FEB-2003 (Rel. 41, Last annotation update)
DE Protease Do-like 1, chloroplast precursor (EC 3.4.21.-).
GN DegP OR DEGPR OR A3G27525 OR K16W12.18.
OS Arabidopsis thaliana (Mouse-ear cress).
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; Rosidae;
OC eucosids II; Brassicales; Brassicaceae; Arabidopsis.
OX NCBI_Taxid=3702;
RN [1]
RP SEQUENCE FROM N.A., AND CHARACTERIZATION.
RX MEDLINE=98175982; PubMed=9507020;
RA Itzhaki H., Naveh L., Lindahl M., Cook M., Adam Z.;
*Identification and characterization of DegP, a serine protease
RT associated with the luminal side of the thylakoid membrane.";
RL J. Biol. Chem. 273:7094-7098(1998).
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=cv. Columbia;

RX MEDLINE:20363099; PubMed:10907853;
 RA Kaneko T., Katoh T., Sato S., Nakamura A., Asamizu E., Tabata S.:
 RT "Structural analysis of Arabidopsis thaliana chromosome 3. II.
 RT Sequence features of the 4,251,695 bp regions covered by 90 Pl. TAC
 RT and BAC clones.";
 RL DNA Res. 7:217-221(2000).
 RN [3]
 RP SEQUENCE OF 104-118.
 RC STRAIN:cv. Columbia;
 RA Kieselbach T., Bystedt M., Schroeder W.P.;
 RL Submitted (JUL-2000) to the SWISS-PROT data bank.
 CC !- FUNCTION: SERINE PROTEASE THAT IS REQUIRED AT HIGH TEMPERATURE.
 CC MAY BE INVOLVED IN THE DEGRADATION OF DAMAGED PROTEINS. IN VIVO.
 CC CAN DEGRADE BETA-CASEIN.
 CC !- ENZYME REGULATION: INHIBITED BY PHENYL METHYLSULFONYL FLUORIDE AND
 CC O-PHENANTHROLINE.
 CC !- SUBCELLULAR LOCATION: BOUND TO LUMINAL SIDE OF THE THYLAKOID
 CC MEMBRANE.
 CC !- INDUCTION: By heat shock.
 CC !- SIMILARITY: BELONGS TO PEPTIDASE FAMILY S2C.
 CC !- SIMILARITY: Contains 1 PDZ/DHR domain.
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 DR EMBL: AF028842; AAC39436.1;
 DR EMBL: AP000371; BAB02539.1;
 DR EMBL: AP001302; BAB02539.1; JOINED.
 DR MEROPS: S01.279;
 DR InterPro: IPR001478; PDZ;
 DR InterPro: IPR001340; Protease2C.
 DR InterPro: IPR001254; Ser_protease_Try.
 DR Pfam: PF00595; PDZ; 1;
 DR PRINTS: PRO0834; PROTEASES2C.
 DR SMART: SM00228; PDZ; 1;
 DR PROSITE: PS50.06; PDZ; 1;
 DR Hydrolase: Serine protease; Transit peptide; Chloroplast; Thylakoid.
 KW TRANSIT ?
 FT TRANSIT 1 ?
 FT CHAIN 104 437
 FT DOMAIN 152 321
 FT DOMAIN 324 421
 FT ACT_SITE 271 171
 FT ACT_SITE 201 202
 FT ACT_SITE 280 280
 FT CONFLICT 12 23
 FT CONFLICT 36 36
 FT CONFLICT 54 54
 FT CONFLICT 60 60
 FT CONFLICT 64 64
 FT CONFLICT 68 69
 FT CONFLICT 355 355
 FT CONFLICT 381 381
 FT CONFLICT 416 416
 SEQUENCE 437 AA: 46213 MW: 149781AB35F5F2A4 CRC64:
 Query Match 9.4%; Score 80.5; DB 1; Length 437;
 Best Local Similarity 25.6%; Pred No. 2.7;
 Matches 44; Conservative 18; Mismatches 55; Indels 55; Gaps 7;
 QY 56 VYHGAGRTIASPKGPIQMY-----TNVQELVGV-----FA 88
 Db 150 VPGSGSGFVMDKQCH:VTNYHVRGASD:RVTLA:ITFDKAVGVGFDKQDVAVLIDA 209
 QY 89 PGGRSLTPTCTCGSSDLYLV-----TRIADYLPVRRERGRGSLSPRT 183

DB 210 PK--NKLRPPIPVGVSADLLVGQKVFALGNPFGLDHTLTITVISGLRREIS--SAATGRPI 265
 QY 134 SYL-----KSGSGGPLLCPAGHVGIPFAAVCTRGVAKAVDF-LPVESL 176
 Db 266 QDVITDAAINPGNSGGLDSSGLIGINTAIYSPGASGSGVGFIPVDIV 317
 RESULT 14
 CAC3_BOVIN
 ID CAC3_BOVIN STANDARD; PRT; 253 AA.
 AC PC5805;
 DT 01-NOV-1988 (Rel. 09, Created)
 DT 15-DEC-1998 (Rel. 37, Last sequence update)
 DT 15-SEP-2003 (Rel. 42, Last annotation update)
 DE Proproteinase F precursor (Procarboxypeptidase A complex component
 DE 11) (Procarboxypeptidase A-S6 subunit III) (PROCPA-S6 III).
 CS Bos taurus (Bovine).
 CC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 CC Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
 CC Bovidae; Bovinae; Bos.
 CX NCBI TaxID=9913;
 RN [1]
 RP SEQUENCE OF 1-25.
 RX MEDLINE:91099520; PubMed:2269366;
 RA Pascua R., Vendrell J., Aviles F.X., Bonicel J., Wicker C.,
 RA Puigserver A.;
 RT "Autolysis of proproteinase F in bovine procarboxypeptidase A ternary
 RT complex gives rise to subunit III.";
 RL FEMS Lett. 277:37-41(1990).
 RN [2]
 RP SEQUENCE OF 14-253, AND DISULFIDE BONDS.
 RX MEDLINE:86220198; PubMed:3519215;
 RA Venot N., Sclaky M., Puigserver A., Desnuelle P., Laurent G.;
 RT "Amino acid sequence and disulfide bridges of subunit III, a
 RT defective carboxypeptidase present in the bovine pancreatic 6 S
 RI procarboxypeptidase A complex.";
 RL Eur. J. Biochem. 157:91-99(1986).
 RN [3]
 RP X-RAY CRYSTALLOGRAPHY (1.7 ANGSTROMS).
 RX MEDLINE:94222022; PubMed:8168476;
 RA Pignol D., Gaboriaud C., Michon I., Kerfelec B., Chapus C.,
 RA Fontecilla-Camps J.C.;
 RT "Crystal structure of bovine procarboxypeptidase A-S6 subunit III, a
 RT highly structured truncated zymogen E.";
 RL EMBO J. 13:1763-1771(1994).
 CC !- FUNCTION: DEFECTIVE ELASTASE-LIKE SERINE PROTEASE. DOES NOT SEEM
 CC TO HAVE A PROTEASE ACTIVITY. ITS LIKELY FUNCTION IS TO PROTECT
 CC PROCARBOXYPEPTIDASE A AGAINST DENATURATION IN THE ACIDIC
 CC ENVIRONMENT OF THE RUMINANT DUODENUM.
 CC !- SUBUNIT: HETEROTRIMER OF SUBUNIT III; CARBOXYPEPTIDASE A AND
 CC CHYMOTRYPSINOGEN C.
 CC !- TISSUE SPECIFICITY: Extracellular.
 CC !- SIMILARITY: BELONGS TO PEPTIDASE FAMILY S1.
 DR PDB: IFON; 14-OCT-96.
 DR PDB: IPYT; 27-JAN-97.
 DR MEROPS: S01.983;
 DR InterPro: IPR001314; Chymotrypsin.
 DR InterPro: IPR001254; Ser_protease_Try.
 DR Pfam: PF00089; trypsin; 1.
 DR PRINTS: PR00722; CHYMOTRYPSIN.
 DR SMART: SM00020; Tryp_Spc; 1.
 DR PROSITE: PS0240; TRYPSIN_DOM; 1.
 DR PROSITE: PS00134; TRYPSIN_HIS; 1.
 DR PROSITE: PS00135; TRYPSIN_SER; 1.
 KW Serine protease homolog; Pancreas; Digestion; 3D-structure.
 FT PROPEP 1 11
 FT CHAIN 12 253
 FT DISULFID 41 57
 FT DISULFID 100 103
 FT DISULFID 140 206
 FT DISULFID 171 187
 FT DISULFID 196 227

A:Title: The Taiwanese hepatitis C virus genome: sequence determination and mapping the
A:Reference number: A40244; MUID:92230206; PMID:1314449

A:Accession: A40244

A:Molecule type: genomic RNA

A:Residues: 1-3010 <CHE>

A:Cross-references: GB:M44754

C:Superfamily: hepatitis C virus genome polyprotein

C:Keywords: ATP; capsid protein; envelope protein; glycoprotein; hydrolase; nonstructural

F:1-115/Product: capsid protein C #status predicted <GPC>

F:116-191/Product: envelope protein M #status predicted <EPM>

F:192-389/Product: major envelope protein E #status predicted <MEP>

F:390-729/Product: nonstructural protein NS1 #status predicted <NS1>

F:730-1006/Product: nonstructural protein NS2 #status predicted <NS2>

F:1007-1615/Product: hepatitis virus #status predicted <NS3>

F:1230-1237/Region: nucleotide-binding motif A (P-loop)

F:1312-1317/Region: nucleotide-binding motif B

F:1316-1319/Region: DEXH motif

F:1516-1862/Product: nonstructural protein NS4 #status predicted <NS4>

F:1863-2013/Product: nonstructural protein NS4b #status predicted <NS4b>

F:2014-3010/Product: nonstructural protein NS5 #status predicted <NS5>

F:196,209,234,250,305,325,417,423,430,448,532,540,556,576,623,645,1213,1255,2041,2077

Query Match 93.6%; Score 892; DB 1; Length 3010;

Best Local Similarity 90.1%; Pred. No. 4,1e-76;

Matches 164; Conservative 12; Mismatches 6; Indels 0; Gaps 0;

QY 1 MAPITAYAOQTGRLGCGITISLTGRDKNOVEGEVOIVSTAAQTFLATCINGVCWTVYHGA 60

DB 1026 LAPITAYAOQTGRLGCGITISLTGRDKNOVEGEVOIVSTAAQTFLATCINGVCWTVYHGA 1085

QY 61 GTRTIASPKGPVIOMYINVDKLVGWPAPQGSRSLSIPTCGSSDLYLVIRHADVIPVRRR 120

DB 1086 GSKTLAAPKGPITOMYINVDKLVGWPAPQGSRSLSIPTCGSSDLYLVIRHADVIPVRRR 1145

QY 121 GDSRGSLLSPRPISYLGKSSGGLPCPAGHAGVIFRAAVCTIRGVAKAVDFIPVESLETTM 180

DB 1146 GDSRGSLLSPRPISYLGKSSGGLPCPAGHAGVIFRAAVCTIRGVAKAVDFIPVESMETTM 1205

QY 181 RS 182

DB 1206 RS 1207

RESULT 5

GNMVCJ

genome polyprotein - hepatitis C virus

N:Contains: capsid protein C; envelope protein M; hepatitis virus (EC 3.4.21.98) (nonstructu

protein NS4a; nonstructural protein NS4b; nonstructural protein NS5

C:Species: hepatitis C virus

C:Date: 31-Mar-1992 #sequence_revision 31-Mar-1992 #text_change 19-Jan-2001

C:Accession: A38465

R:Takamizawa, A.; Wori, C.; Fuke, I.; Manabe, S.; Murakami, S.; Fujita, J.; Onishi, E.;

J. Virol. 65, 1105-1113, 1991

A:Title: Structure and organization of the hepatitis C virus genome isolated from human

A:Reference number: A38465; MUID:91140698; PMID:1847440

A:Accession: A38465

A:Molecule type: genomic RNA

A:Residues: 1-3010 <TA>

A:Cross-references: EMBL:M58335; NID:g329770; PID:AAA72945.1; PID:g329771

C:Superfamily: hepatitis C virus genome polyprotein

C:Keywords: ATP; capsid protein; envelope protein; glycoprotein; hydrolase; nonstructural

F:1-115/Product: capsid protein C #status predicted <GPC>

F:116-191/Product: envelope protein M #status predicted <EPM>

F:192-389/Product: major envelope protein E #status predicted <MEP>

F:390-729/Product: nonstructural protein NS1 #status predicted <NS1>

F:730-1006/Product: nonstructural protein NS2 #status predicted <NS2>

F:1007-1615/Product: hepatitis virus #status predicted <NS3>

F:1230-1237/Region: nucleotide-binding motif A (P-loop)

F:1312-1317/Region: nucleotide-binding motif B

F:1316-1319/Region: DEXH motif

F:1516-1862/Product: nonstructural protein NS4 #status predicted <NS4>

F:1863-2013/Product: nonstructural protein NS4b #status predicted <NS4b>

F:2014-3010/Product: nonstructural protein NS5 #status predicted <NS5>

F:196,209,234,250,305,325,417,423,430,448,532,540,556,576,623,645,1213,1255,2041,2077

Query Match 93.5%; Score 891; DB 1; Length 3010;

Best Local Similarity 89.0%; Pred. No. 5,1e-76;

Matches 162; Conservative 15; Mismatches 5; Indels 0; Gaps 0;

QY 1 MAPITAYAOQTGRLGCGITISLTGRDKNOVEGEVOIVSTAAQTFLATCINGVCWTVYHGA 60

DB 1026 LAPITAYAOQTGRLGCGITISLTGRDKNOVEGEVOIVSTAAQTFLATCINGVCWTVYHGA 1085

QY 61 GTRTIASPKGPVIOMYINVDKLVGWPAPQGSRSLSIPTCGSSDLYLVIRHADVIPVRRR 120

DB 1086 GSKTLAAPKGPITOMYINVDKLVGWPAPQGSRSLSIPTCGSSDLYLVIRHADVIPVRRR 1145

QY 121 GDSRGSLLSPRPISYLGKSSGGLPCPAGHAGVIFRAAVCTIRGVAKAVDFIPVESLETTM 180

DB 1146 GDSRGSLLSPRPISYLGKSSGGLPCPAGHAGVIFRAAVCTIRGVAKAVDFIPVESMETTM 1205

QY 181 RS 182

DB 1206 RS 1207

RESULT 6

GNMVCJ

genome polyprotein - hepatitis C virus (strain J)

N:Contains: capsid protein C; envelope protein M; major envelope protein E; nonstru

protein NS4a; nonstructural protein NS4b; nonstructural protein NS5

C:Species: hepatitis C virus

C:Date: 30-Jun-1992 #sequence_revision 30-Jun-1992 #text_change 19-Jan-2001

C:Accession: A39253; PS0086

R:Kato, N.; Hijikata, M.; Ootsuyama, Y.; Nakagawa, M.; Ohkoshi, S.; Sugimura, T.;

Proc. Natl. Acad. Sci. U.S.A. 87, 9524-9528, 1990

A:Title: Molecular cloning of the human hepatitis C virus genome from Japanese pat.

A:Reference number: A39253; MUID:91088550; PMID:2175903

A:Accession: A39253

A:Molecule type: genomic RNA

A:Residues: 1-3010 <KA7>

A:Cross-references: GB:D90208; NID:g221610; PID:BAAL4233.1; PID:g221611

R:Kato, N.; Ohkoshi, S.; Shimotohno, K.

Proc. Jpn. Acad. 65B, 219-223, 1989

A:Title: Japanese isolates of the non-A, non-B hepatitis viral genome show sequen

A:Reference number: PS0085

A:Accession: PS0086

A:Molecule type: genomic RNA

A:Residues: 2650-2707 <KA2>

A:Experimental source: Japanese isolate

C:Comment: The cleavage sites of this polyprotein have not been determined.

C:Superfamily: hepatitis C virus genome polyprotein

C:Keywords: ATP; glycoprotein; hydrolase; nucleotide binding; P-loop; polyprotein;

F:2-115/Product: capsid protein C #status predicted <GPC>

F:116-191/Product: major envelope protein M #status predicted <EPM>

F:192-389/Product: nonstructural protein NS1 #status predicted <NS1>

F:390-729/Product: nonstructural protein NS2 #status predicted <NS2>

F:730-1006/Product: nonstructural protein NS2 #status predicted <NS2>

F:1007-1615/Product: hepatitis virus #status predicted <NS3>

F:1230-1237/Region: nucleotide-binding motif A (P-loop)

F:1312-1317/Region: nucleotide-binding motif B

F:1316-1319/Region: DEXH motif

F:1516-1862/Product: nonstructural protein NS4 #status predicted <NS4>

F:1863-2013/Product: nonstructural protein NS4b #status predicted <NS4b>

F:2014-3010/Product: nonstructural protein NS5 #status predicted <NS5>

F:196,209,234,250,305,325,417,423,430,448,532,556,576,623,645,1213,1255,2041,2077,

Query Match 93.1%; Score 887; DB 1; Length 3010;

Best Local Similarity 87.9%; Pred. No. 1,2e-75;

Matches 160; Conservative 17; Mismatches 5; Indels 0; Gaps 0;

QY 1 MAPITAYAOQTGRLGCGITISLTGRDKNOVEGEVOIVSTAAQTFLATCINGVCWTVYHGA 60

DB 1026 LAPITAYAOQTGRLGCGITISLTGRDKNOVEGEVOIVSTAAQTFLATCINGVCWTVYHGA 1085

QY 61 GTRTIASPKGPVIOMYINVDKLVGWPAPQGSRSLSIPTCGSSDLYLVIRHADVIPVRRR 120

```
1086 GSKTAGKPGITQYNTVNDQVLGVHPAPPCASMPCTCGSSDLVLTIRHADVVPVRR 1145
QY 121 GDSRGSLLSPRPISYLKSSGGPLCPAGHAGVIFRAAVCTRGVAKAVDFIPVESLETTM 180
DB 1146 GDSRGSLLSPRPISYLKSSGGPLCPAGHAGVIFRAAVCTRGVAKAVDFIPVESMETTM 1205
QY 181 RS 182
DB 1206 RS 1207

RESULT 7
genome polyprotein - hepatitis C virus (strain J1)
N:Contains: capsid protein C; envelope protein M; hepatitis C virus (nonstructural protein NS4b; nonstructural protein NS5)
C:Species: hepatitis C virus
C:Date: 19-May-2000 #sequence_revision 19-May-2000 #text_change 19-Jan-2001
C:Accession: A45573
R:Tanaka, T.; Kato, N.; Nakagawa, M.; Ootsuyama, Y.; Cho, M.J.; Nakazawa, T.; Hijikata, R.
A:Title: Molecular cloning of hepatitis C virus genome from a single Japanese carrier
A:Reference number: A45573; MUID:92295714; PMID:1335627
A:Accession: A45573
A>Status: preliminary
A:Molecule type: DNA
A:Residues: 1-3010 <FAN>
A:Cross-references: GB:D01171; NID:g221612; PIDN:BAA01943.1; PID:g221613
A:Experimental source: HCV-JT
A>Note: sequence extracted from NCBI backbone (NCBI:106206, NCBI:106207)
C:Superfamily: hepatitis C virus genome polyprotein
C:Keywords: ATP; glycoprotein; hydrolase; nucleotide binding; P-loop; polyprotein; serine
F:2-115/Product: capsid protein C #status predicted <CPC>
F:116-191/Product: envelope protein M #status predicted <EPM>
F:192-389/Product: major envelope protein E #status predicted <MEP>
F:390-729/Product: nonstructural protein NS1 #status predicted <NS1>
F:730-1006/Product: nonstructural protein NS2 #status predicted <NS2>
F:1007-1615/Product: hepatitis C virus predicted <NS3>
F:1230-1237/Region: nucleotide-binding motif A (P-loop)
F:1312-1317/Region: nucleotide-binding motif B
F:1316-1319/Region: DEXH motif
F:1616-1862/Product: nonstructural protein NS4a #status predicted <N4A>
F:1863-2013/Product: nonstructural protein NS4b #status predicted <N4B>
F:2014-3010/Product: nonstructural protein NS5 #status predicted <NS5>

Query Match 92.8%; Score 884; DB 1; Length 3010;
Best Local Similarity 88.5%; Pred. No. 2.3e-75;
Matches 161; Conservative 15; Mismatches 6; Indels 0; Gaps 0;

QY 1 MAPITAYAOQTRGLGCIITSLTGDRKNQVGEVQIVSTAAQIFATCINXGVCTVYHGA 60
DB 1026 LAPITAYAOQTRGLGCIITSLTGDRKNQVGEVQIVSTAAQIFATCINXGVCTVYHGA 1085
QY 61 GTRITASPKGPVQMYINVDKLVGWAPQSRSLTCTCGSSDLVLTIRHADVVPVRR 120
DB 1086 GSKTAGKPGITQYNTVNDQVLGVHPAPPCASMPCTCGSSDLVLTIRHADVVPVRR 1145
QY 121 GDSRGSLLSPRPISYLKSSGGPLCPAGHAGVIFRAAVCTRGVAKAVDFIPVESLETTM 180
DB 1146 GDSRGSLLSPRPISYLKSSGGPLCPAGHAGVIFRAAVCTRGVAKAVDFIPVESMETTM 1205
QY 181 RS 182
DB 1206 RS 1207

RESULT 8
S18030
genome polyprotein - hepatitis C virus (isolate J1)
N:Contains: capsid protein C; envelope protein M; hepatitis C virus (nonstructural protein NS4b; nonstructural protein NS5)
C:Species: hepatitis C virus
C:Date: 19-May-2000 #sequence_revision 19-May-2000 #text_change 19-Jan-2001
C:Accession: J18030
R:Chamberlain, R.W.; Adams, N.J.; Taylor, L.A.; Simmonds, P.; Elliott, R.M.
A:Title: The complete coding sequence of hepatitis C virus genotype 5a, the predominant
A:Reference number: J18030; MUID:97366593; PMID:9223423
```

```
A:Variety: isolate J1
C:Date: 19-May-2000 #sequence_revision 19-May-2000 #text_change 23-Mar-2001
C:Accession: S18030; S33570; A48332; S18029
R:Honda, M.; Kaneko, S.; Masashi, U.; Kobayashi, K.; Murakami, S.
A:Title: The complete coding sequence of hepatitis C virus genotype 5a, the predominant
A:Reference number: S18028
A:Accession: S18030
A:Molecule type: genomic RNA
A:Residues: 1-3010 <HON>
A:Cross-references: EMBL:X61596; NID:g59478; PIDN:CAA4793.1; PID:g59479
A:Experimental source: isolate J1 from an individual
R:Honda, M.; Kaneko, S.; Ueno, M.; Kobayashi, K.; Murakami, S.
A:Title: Sequence analysis of putative structural regions of hepatitis C virus isolate
A:Reference number: A48332; MUID:9119270; PMID:8380322
A:Accession: S33570
A:Residues: 1-547, 'T', 549-621, 'V', 623-624, 'S', 626-652, 'DL', 655-761, 'T', 763-782 <HOW>
A:Cross-references: EMBL:X61591
A>Note: this sequence is inconsistent with the nucleotide translation
as 3TP, and TTC for residue 771 as Ser
A>Note: sequence extracted from NCBI backbone (NCBI:121747, NCBI:121748)
C:Superfamily: hepatitis C virus genome polyprotein
C:Keywords: ATP; glycoprotein; hydrolase; nucleotide binding; P-loop; polyprotein;
F:2-115/Product: capsid protein C #status predicted <CPC>
F:116-191/Product: envelope protein M #status predicted <EPM>
F:192-389/Product: major envelope protein E #status predicted <MEP>
F:390-729/Product: nonstructural protein NS1 #status predicted <NS1>
F:730-1006/Product: nonstructural protein NS2 #status predicted <NS2>
F:1007-1615/Product: hepatitis C virus predicted <NS3>
F:1230-1237/Region: nucleotide-binding motif A (P-loop)
F:1312-1317/Region: nucleotide-binding motif B
F:1316-1319/Region: DEXH motif
F:1616-1862/Product: nonstructural protein NS4a #status predicted <N4A>
F:1863-2013/Product: nonstructural protein NS4b #status predicted <N4B>
F:2014-3010/Product: nonstructural protein NS5 #status predicted <NS5>
F:196,209,234,250,305,417,423,448,532,540,556,576,623,643,645/Binding site: carbohydrate

Query Match 90.7%; Score 864; DB 1; Length 3010;
Best Local Similarity 86.8%; Pred. No. 1.9e-73;
Matches 158; Conservative 15; Mismatches 9; Indels 0; Gaps 0;

QY 1 MAPITAYAOQTRGLGCIITSLTGDRKNQVGEVQIVSTAAQIFATCINXGVCTVYHGA 60
DB 1026 LAPITAYAOQTRGLGCIITSLTGDRKNQVGEVQIVSTAAQIFATCINXGVCTVYHGA 1085
QY 61 GTRITASPKGPVQMYINVDKLVGWAPQSRSLTCTCGSSDLVLTIRHADVVPVRR 120
DB 1086 GSKTAGKPGITQYNTVNDQVLGVHPAPPCASMPCTCGSSDLVLTIRHADVVPVRR 1145
QY 121 GDSRGSLLSPRPISYLKSSGGPLCPAGHAGVIFRAAVCTRGVAKAVDFIPVESLETTM 180
DB 1146 GDSRGSLLSPRPISYLKSSGGPLCPAGHAGVIFRAAVCTRGVAKAVDFIPVESMETTM 1205
QY 181 RS 182
DB 1206 RS 1207

RESULT 9
JC5620
genome polyprotein - hepatitis C virus (isolate EUH1480)
N:Contains: capsid protein C; envelope protein M; hepatitis C virus (nonstructural protein NS4b; nonstructural protein NS5)
C:Species: hepatitis C virus
C:Date: 19-May-2000 #sequence_revision 19-May-2000 #text_change 19-Jan-2001
C:Accession: JC5620
R:Chamberlain, R.W.; Adams, N.J.; Taylor, L.A.; Simmonds, P.; Elliott, R.M.
A:Title: The complete coding sequence of hepatitis C virus genotype 5a, the predominant
A:Reference number: JC5620; MUID:97366593; PMID:9223423
```

A:Accession: J00559
A:Molecule type: mRNA
A:Residues: 2678-2729 <KAT>
A:Cross-references: GB:D0562; GB:D90518; NID:g221523; PIDN:BA04148.1; PID:g221523.2
A:Experimental source: genotype 5a, which predominates in South Africa
A:Note: the translation of the nucleotide sequence is not complete in this paper
C:Superfamily: hepatitis C virus genome polyprotein
C:Keywords: ATP; capsid protein; envelope protein; glycoprotein; hydrolase; serine proteinase;
F:115-191/Product: capsid protein C #status predicted <CPC>
F:116-191/Product: capsid protein C #status predicted <CPC>
F:192-389/Product: major envelope protein E #status predicted <EPM>
F:192-389/Product: major envelope protein E #status predicted <EPM>
F:192-389/Product: major envelope protein E #status predicted <EPM>
F:390-733/Product: nonstructural protein NS1 #status predicted <NS1>
F:734-1210/Product: nonstructural protein NS2 #status predicted <NS2>
F:1011-1619/Product: hepatitis C virus genome polyprotein
F:1234-1242/Region: nucleotide-binding motif A (P-loop)
F:1234-1242/Region: nucleotide-binding motif B
F:1316-1321/Region: DEXH motif
F:1320-1323/Region: DEXH motif
F:1520-1866/Product: nonstructural protein NS4a #status predicted <NS4a>
F:1520-1866/Product: nonstructural protein NS4b #status predicted <NS4b>
F:1567-2017/Product: nonstructural protein NS5 #status predicted <NS5>
F:2018-3033/Product: nonstructural protein NS5 #status predicted <NS5>
F:196,299,234,305,417,423,430,448,477,534,542,558,578,627,649,1091,1217,1259,1259.2

Query Match 84.4%; Score 604; DB 1; Length 3014;
Best Local Similarity 79.1%; Pred. No. 9.5e-68;
Matches 144; Conservative 22; Mismatches 16; Indels 0; Gaps 0;

QY 1 MAPITAYAAQCTGGLGCIITSLTGDRKNOVEGEVQIVSTAQTFLAICINGVCWTIVYHGA 60
Db 1027 LAPITAYAAQCTGAGVATSLTGDRKNEAGEVCFSLTATQFELATCINGVWMTLFRHA 1484
QY 61 GTRTASPKGPVYOMTYNDKLVGMWPAFGSGRLTCTGCGSSDLYLVIRHADVTPVRRR 120
Db 1087 GSKLAGKPGPVOMTYNDKLVGMWPAFGSGRLTCTGCGSSDLYLVIRHADVTPVRRR 1146
QY 121 GDSRGLSPRISYILKSGSGGLPCLPAGHAGVIFRAAVCTRGVAKAVDFIPVESLETTM 180
Db 1247 GDSRGLSPRISYILKSGSGGLPCLPAGHAGVIFRAAVCTRGVAKAVDFIPVESLETTM 180
QY 161 RS 162
Db 1207 RS 1208

RESULT 10
GENWJ8
genome polyprotein - hepatitis C virus (strain HC-J8)
N:Contains: capsid protein C; envelope protein M; hepatitis C virus genome polyprotein
C:Keywords: ATP; capsid protein; envelope protein; glycoprotein; hydrolase; serine proteinase;
F:115-191/Product: capsid protein C #status predicted <CPC>
F:116-191/Product: capsid protein C #status predicted <CPC>
F:192-389/Product: major envelope protein E #status predicted <EPM>
F:192-389/Product: major envelope protein E #status predicted <EPM>
F:390-733/Product: nonstructural protein NS1 #status predicted <NS1>
F:734-1210/Product: nonstructural protein NS2 #status predicted <NS2>
F:1011-1619/Product: hepatitis C virus genome polyprotein
F:1234-1242/Region: nucleotide-binding motif A (P-loop)
F:1234-1242/Region: nucleotide-binding motif B
F:1316-1321/Region: DEXH motif
F:1320-1323/Region: DEXH motif
F:1520-1866/Product: nonstructural protein NS4a #status predicted <NS4a>
F:1520-1866/Product: nonstructural protein NS4b #status predicted <NS4b>
F:1567-2017/Product: nonstructural protein NS5 #status predicted <NS5>
F:2018-3033/Product: nonstructural protein NS5 #status predicted <NS5>
F:196,299,234,305,417,423,430,448,477,534,542,558,578,627,649,1091,1217,1259.2

Query Match 84.4%; Score 604; DB 1; Length 3014;
Best Local Similarity 79.1%; Pred. No. 9.5e-68;
Matches 144; Conservative 22; Mismatches 16; Indels 0; Gaps 0;

QY 1 MAPITAYAAQCTGGLGCIITSLTGDRKNOVEGEVQIVSTAQTFLAICINGVCWTIVYHGA 60
Db 1027 LAPITAYAAQCTGAGVATSLTGDRKNEAGEVCFSLTATQFELATCINGVWMTLFRHA 1484
QY 61 GTRTASPKGPVYOMTYNDKLVGMWPAFGSGRLTCTGCGSSDLYLVIRHADVTPVRRR 120
Db 1087 GSKLAGKPGPVOMTYNDKLVGMWPAFGSGRLTCTGCGSSDLYLVIRHADVTPVRRR 1146
QY 121 GDSRGLSPRISYILKSGSGGLPCLPAGHAGVIFRAAVCTRGVAKAVDFIPVESLETTM 180
Db 1247 GDSRGLSPRISYILKSGSGGLPCLPAGHAGVIFRAAVCTRGVAKAVDFIPVESLETTM 180
QY 161 RS 162
Db 1207 RS 1208

RESULT 10
GENWJ8
genome polyprotein - hepatitis C virus (strain HC-J8)
N:Contains: capsid protein C; envelope protein M; hepatitis C virus genome polyprotein
C:Keywords: ATP; capsid protein; envelope protein; glycoprotein; hydrolase; serine proteinase;
F:115-191/Product: capsid protein C #status predicted <CPC>
F:116-191/Product: capsid protein C #status predicted <CPC>
F:192-389/Product: major envelope protein E #status predicted <EPM>
F:192-389/Product: major envelope protein E #status predicted <EPM>
F:390-733/Product: nonstructural protein NS1 #status predicted <NS1>
F:734-1210/Product: nonstructural protein NS2 #status predicted <NS2>
F:1011-1619/Product: hepatitis C virus genome polyprotein
F:1234-1242/Region: nucleotide-binding motif A (P-loop)
F:1234-1242/Region: nucleotide-binding motif B
F:1316-1321/Region: DEXH motif
F:1320-1323/Region: DEXH motif
F:1520-1866/Product: nonstructural protein NS4a #status predicted <NS4a>
F:1520-1866/Product: nonstructural protein NS4b #status predicted <NS4b>
F:1567-2017/Product: nonstructural protein NS5 #status predicted <NS5>
F:2018-3033/Product: nonstructural protein NS5 #status predicted <NS5>
F:196,299,234,305,417,423,430,448,477,534,542,558,578,627,649,1091,1217,1259.2

Query Match 74.7% Score 712; DB 2; Length 1043;
Best Local Similarity 54.8%; Pred. No. 54e-59;
Matches 127; Conservative 29; Mismatches 26; Indels 0; Gaps 0;
QY 1 KAPITAYAOQTGRLGCIITSLTGRDKNQVEGEVQIVSTAAQTFLATCINGVGVVYHGA 60
DB 2030 KAPITAYAOQTGRLGCIITSLTGRDKNQVEGEVQIVSTAAQTFLATCINGVGVVYHGA 1099
QY 61 CTRTIASPKGPVIONYTNVDKLVGWPAPQGSRLTPTCTGSSDLVLTTRHADVPVRRG 120
DB 1090 GNTIAGSRGPVIONYTNVDKLVGWPAPQGSRLTPTCTGSSDLVLTTRHADVPVRRG 1149
QY 121 GDSRGLSPRPISYLGKSSGGPLCPAGHAGVIFRAAVCTRGVAKAVDFIPVESLET 180
DB 1150 GDRKGA-LSPRLSTLKGSSGGPLCPRHAGVGVFRAAVCTRGVAKAVDFIPVESLET 1269
QY 181 RS 182
DB 1210 RS 1211
RESULT 12
T08841
Polyprotein - douroucouli hepatitis GB virus A
C:Species: douroucouli hepatitis GB virus A
C:Date: 20-Sep-1999 #sequence_revision 20-Sep-1999 #text_change 17-Nov-2000
C:Accession: T08841
R:Erker, J.C.; Desai, S.M.; Leary, T.P.; Chalmers, M.L.; Montes, C.C.; Mushahwar, I.K.
J. Gen. Virol. 79, 41-45, 1998
A:Title: Genomic analysis of two GB virus A variants isolated from captive monkeys.
A:Reference number: Z16486; MUID:98120818; PMID:9460920
A:Accession: T08841
A:Molecule type: mRNA
A:Status: translated from GB/EMBL/DBJ
A:Residues: 1-3005 <ERK>
A:Cross-references: EMBL:AF023425; NID:g2828599; PIDN:AAC40502.1; PID:g2828630
A:Superfamily: hepatitis C virus genome polyprotein
C:Keywords: polyprotein
Query Match 28.1% Score 267.5; DB 2; Length 3005;
Best Local Similarity 33.1%; Pred. No. 1e-16;
Matches 60; Conservative 32; Mismatches 75; Indels 11; Gaps 4;
QY 2 APTIAYAOQTGRLGCIITSLTGRDKNQVEGEVQIVSTAAQTFLATCINGVGVVYHGA 61
DB 579 APVVV-WORGLGPFSSVVKTSMEGRDEREHEGSIIVLTSTIRSMGICVNGVMTTFHGSN 1037
QY 52 IRTIASPKGPVIONYTNVDKLVGWPAPQGSRLTPTCTGSSDLVLTTRHADVPVRRG 121
DB 1038 ARTIAGVGVPCNCRWSPSDVAVYPTPSGASCTHEPKGCTQSVWCTRN--DGAICGR 1095
QY 122 DSRGSLSPRPISYLGKSSGGPLCPAGHAGVIFRAAVCTRGVAKAVDFIPVE 174
DB 1096 SKVLVDLPTTISDFRSGSSGSLCDESHVGVGM-VSVLHRGVKATGVAVRVKFWH:LFK 1154
QY 175 S 175
DB 1155 S 1155
RESULT 13
T08839
polyprotein - marmoset hepatitis GB virus A
C:Species: marmoset hepatitis GB virus A
C:Date: 23-Jul-1999 #sequence_revision 23-Jul-1999 #text_change 17-Nov-2000
C:Accession: T08839
R:Erker, J.C.; Desai, S.M.; Leary, T.P.; Chalmers, M.L.; Montes, C.C.; Mushahwar, I.K.
J. Gen. Virol. 79, 41-45, 1998
A:Title: Genomic analysis of two GB virus A variants isolated from captive monkeys.
A:Reference number: Z16486; MUID:98120818; PMID:9460920
A:Accession: T08839
A:Status: translated from GB/EMBL/DBJ
A:Molecule type: genomic RNA

A:Residues: 1-2970 <ERK>
A:Cross-references: EMBL:AF023424; NID:g2828597; PIDN:AAC40501.1; PID:g2828598
A:Superfamily: hepatitis C virus genome polyprotein
C:Keywords: polyprotein
Query Match 26.8% Score 255.5; DB 2; Length 2970;
Best Local Similarity 30.1%; Pred. No. 1.4e-15;
Matches 59; Conservative 36; Mismatches 66; Indels 33; Gaps 6;
QY 2 APTIAYAOQTGRLGCIITSLTGRDKNQVEGEVQIVSTAAQTFLATCINGVGVVYHGA 61
DB 970 APVVV-HHGKGFVGVKISMTGDETHGVNVVVLGISTRSMTGTCVNVMTTHGSN 1028
QY 62 IRTIASPKGPVIONYTNVDKLVGWPAPQGSRLTPTCTGSSDLVLTTRHADVPVRRG 121
DB 1029 ARTIAGVGVPCNCRWSPSDVAVYPTPSGASCTHEPKGCTQSVWCTRN--DGAICGR 1095
QY 122 DSRGSLSPRPISYLGKSSGGPLCPAGHAGVIFRAAVCTRGVAKAVDFIPVESLET 179
DB 1078 D--GALCHCTGRTVELDPAELCDFRSGSPILCDGHAAGML-ISVLRGSRVTGIR 1134
QY 164 VAKAVDFIPVESLET 179
DB 1135 YTKPWETLPREATHY 1150
RESULT 14
H83144
probable aromatic acid decarboxylase PA4019 [imported] - Pseudomonas aeruginosa (str)
C:Species: Pseudomonas aeruginosa
C:Date: 15-Sep-2000 #sequence_revision 15-Sep-2000 #text_change 31-Dec-2000
C:Accession: H83144
R:Stover, C.K.; Pham, X.Q.; Erwin, A.L.; Mizoguchi, S.D.; Warren, P.; Hickey, M.J.
adman, S.; Yuan, Y.; Brody, L.L.; Coulter, S.N.; Folger, K.R.; Kas, A.; Latbig, K.;
Lory, S.; Olson, M.V.
Nature 406, 959-964, 2000
A:Title: Complete genome sequence of Pseudomonas aeruginosa PA01, an opportunistic
A:Reference number: A82950; MUID:20437337; PMID:10964043
A:Accession: H83144
A:Molecule type: DNA
A:Status: preliminary
A:Residues: 1-209 <STO>
A:Cross-references: GB:AE004816; GB:AE004091; NID:g9950200; PIDN:AAG07406.1; GSPDS:
A:Experimental source: strain PA01
C:Genetics:
A:Gene: PA4019
C:Superfamily: dedF protein
Query Match 9.1% Score 87; DB 2; Length 205;
Best Local Similarity 26.2%; Pred. No. 0.69;
Matches 55; Conservative 18; Mismatches 61; Indels 76; Gaps 12;
QY 8 AQQTGRLGCIITSLTGRDKNQVEGEVQIVSTAAQTFLATCINGVGVVYHGA 66
DB 17 AQYGLRLDCLV-----QGEVHFLISKRAQLVMAI-----ETDVA 53
QY 67 SPKGP-----VIQMYTNVDKLVGWPAPQGSRLTPTCTGSSDL 105
DB 54 LPAPQAMQAFLEYCGAAGQIRVFGND-----WAPASGSSAPNMMVICPSTGIL 108
QY 106 -----YLVTRHADVPVRRGDSRGLSPRPIS-----YLGKSSGGPLCPA 148
DB 109 SAVATGACNNIFRAADVALKER-----RPLVLPREAPFSSIHLENMLKLSNLAJAVILPA 164
QY 149 GHAVGIFRAAVCTRGVAKAVDFIPVESLET 178
DB 165 ---APGFYHQ---POSVEDLYDFVVARILNT 189
RESULT 15
I39383
angio-associated migratory cell protein - human
C:Species: Homo sapiens (man)

C:Date: 06-Sep-1996 #sequence_revision 06-Sep-1996 #text_change 21-Jul-2000
 C:Accession: 139383
 R:Beckner, M.E.; Krutzsch, H.C.; Stracke, M.L.; Williams, S.T.; Gallardo, J.A.; Liotta, Cancer Res. 55, 2140-2149, 1995
 A:Title: Identification of a new immunoglobulin superfamily protein expressed in blood v
 A:Reference number: 139383; PMID:95262124; PMID:7743515
 A:Accession: 139383
 A>Status: preliminary; translated from GR/EMBL/DDBJ
 A:Molecule type: mRNA
 A:Residues: 1-452 <RES>
 A:Cross-references: GR:M95627; NID:q870602; PID:AAA68889.1; PID:q870803
 C:Genetics:
 A:Gene: GDR:AAMP
 A:Cross-references: GDB:4573393
 A:Map position: 14q32.1-14q32.1
 C:Superfamily: unassigned WD repeat proteins; WD repeat homology
 F:148-181/Domain: WD repeat homology <WD1>
 F:414-447/Domain: WD repeat homology <WD2>

Query Match 8.64; Score 82; DB 2; Length 452;
 Best Local Similarity 25.3%; Pred. No. 5;
 Matches 42; Conservative 13; Mismatches 47; Indels 44; Gaps 9;
 QY 54 WTVEGCASTETIASPKGPVIGWYTNVDKILYGVWPAQGSRL-----TQCTGSSDLYLV 108
 DB 197 KWEWH-----PRAPVLLAST-ADGNTWVKVPNGCK-FOGPNCFATCGK----- 240
 QY 109 TRHADVIPVRRR---GDSRGS-----LSRPPISTYKQSSG--GPILCFA----- 148
 DB 241 -----VLEPGKRAVGYEETGIRINDELKGGSPHVKETEGHGPGPLTCVAANQDSLLI 295
 QY 149 -----CHAVGIER-----AAVCTRGVAKAVDFIIVESL 179
 DB 296 GSVDQAKLVNATTKGVVPRPRTVASQPSLGGEGESFSNSVESL 341

Search completed: September 27, 2003, 12:22:12
 Job time : 44 secs

GenCore version 5.1.6
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OM protein - nucleic search, using frame_plus_p2n model

Run on: September 27, 2003, 12:28:45 : Search time 2332 Seconds
(without alignments)
1981.817 Million c.c.: updates/sec

Title: US-09-965-594-1
Perfect score: 953
Sequence: 1 MAPITAYAQQRGLLSC-IT.....GVAVADFPVLESLETIMRS 182

Scoring table: BLOSUM62
Xgapop 10.0, Xgapext 0.5
Ygapop 10.0, Ygapext 0.5
Fgapop 6.0, Fgapext 7.0
Delop 6.0, Delext 7.0

Searched: 22781192 seqs, 12152238056 residues

Total number of hits satisfying chosen parameters: 4536274

Minimum DB seq length: 0
Maximum DB seq length: 200000000

Post-processing: Minimum Match 04
Maximum Match 994
Listing first 45 summaries

Command line parameters:
-MODEL=frame-p2n.model -DEV=xih
-O=/cgn2_1/USPTO.spool/US09965594/runat_26092003_164143_18769/aff_query_fastal.1.327
-DB=EST -QFMT=fastap -SUFFIX=first -MINMATCH=9.1 -LOOPEL=0 -DOPEX=0
-UNITS=bits -START=1 -END=1 -MATRIX=blosum62 -TRANS=human40.cgi -LIST=45
-DOCALIGN=200 -THR_SCORE=pct -THR_MAX=99.9 -THR_MIN=0 -ALIGN=15 -MCDE=LOCAL
-OUTFMT=pco -NORM=ext -HEADSIZE=500 -MINLEN=0 -MAXLEN=200000000
-USER=US09965594 -CGEN 1_1_2810 -runat_26092003_164143_18769 -NCPU=6 -ICPU=3
-NO_MMAP -LARGEQUERY -NEG_SCORES=0 -WAIT_DSPBLOCK=100 -LONGLOG
-DEV_TIMEOUT=120 -WARN_TIMEOUT=30 -THREADS=1 -XGAPOP=10 -XGAPEXT=0.5 -Fgapop=6
-Fgapext=7 -YGAPOP=10 -YGAPEXT=9.5 -DELOP=6 -DELEXT=7

Database :

EST :
1: cm_estba1
2: cm_estcum1
3: cm_estcum2
4: cm_estcum3
5: cm_estcum4
6: cm_estcum5
7: cm_estcum6
8: cm_hic1
9: qb_est1
10: qb_est2
11: qb_hic1
12: qb_est3
13: qb_est4
14: qb_est5
15: cm_estcum1
16: cm_estcum2
17: cm_estcum3
18: cm_gss_hum1
19: cm_gss_hum2
20: cm_gss_pln1
21: cm_gss_vrt1
22: cm_gss_fun1
23: cm_gss_fun2
24: cm_gss_mus1
25: cm_gss_mus2
26: cm_gss_pro1
27: cm_gss_pro2
28: cm_gss_phq1
29: cm_gss_vrl1
30: cm_gss_vrl2

29: gb_gss2:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match %	Length	DB	ID	Description
C 1	106	11.1	984	10	BF304699	601888252
C 2	103.5	10.9	1199	13	B0892487	B0892487 AGENCOURT
C 3	101	10.6	515	14	CA023748	CA023748 H247817
C 4	101	10.6	583	12	BM374064	BM374064 Ema03_S0
C 5	99	10.4	515	12	BJ001625	BJ001625 BJO01625
C 6	99	10.4	543	12	BJ024121	BJ024121 BJO24121
C 7	99	10.4	754	12	BJ016176	BJ016176 BJO16176
C 8	98.5	10.3	961	10	BF203316	BF203316 601865914
C 9	98.5	10.3	1031	14	CB950999	CB950999 AGENCOURT
C 10	98.5	10.3	1141	11	AK080545	AK080545 MUS MUSC1
C 11	97.5	10.2	779	10	BF631437	BF631437 HVSMB001
C 12	96.5	10.1	844	11	CNS09DAS	CNS09DAS Single re
C 13	96	10.1	701	10	BF863244	BF863244 963042C02
C 14	95	10.1	546	10	BF182274	BF182274 601804028
C 15	95.5	10.0	901	10	BF307233	BF307233 601891502
C 16	95.5	10.0	958	10	BG420860	BG420860 602452062
C 17	95	10.0	407	9	AW785806	AW785806 117260_MA
C 18	95	10.0	460	14	CB883286	CB883286 H001M02W
C 19	94.5	9.9	539	28	BH349665	BH349665 CH230-65E
C 20	94	9.9	582	14	CB286751	CB286751 CMD45_C08
C 21	94	9.9	1052	10	EG398041	EG398041 602439571
C 22	94	9.9	1263	13	BQ709745	BQ709745 AGENCOURT
C 23	93.5	9.8	736	12	BI768830	BI768830 603057734
C 24	93.5	9.8	938	13	BQ894657	BQ894657 AGENCOURT
C 25	93.5	9.8	993	9	AL555424	AL555424 AL555424
C 26	93	9.8	457	29	CNS02YOL	AL219990 Tetradon
C 27	93	9.8	470	13	BQ758584	BQ758584 Ema07_S0
C 28	93	9.8	1018	12	BQ054587	BQ054587 AGENCOURT
C 29	93	9.8	1291	10	BE622016	BE622016 601440668
C 30	92.5	9.7	655	14	CB888789	CB888789 H009014W
C 31	92.5	9.7	1008	12	BI755608	BI755608 603027112
C 32	92.5	9.7	1411	11	BC020343	BC020343 Homo sap1
C 33	92	9.7	756	14	CD348815	CD348815 UT-M-FYC-
C 34	92	9.7	832	10	BG387051	BG387051 602454749
C 35	92	9.7	871	10	BG178418	BG178418 602330206
C 36	92	9.7	898	29	CNS01VRS	ALL69466 Tetradon
C 37	92	9.7	963	10	BF794182	BF794182 602255566
C 38	92	9.7	1329	13	BQ960995	BQ960995 AGENCOURT
C 39	92	9.7	1540	10	BF180599	BF180599 601808704
C 40	91.5	9.6	422	14	CB763743	CB763743 AMGNNUC:S
C 41	91.5	9.6	539	10	BE757615	BE757615 212104_MA
C 42	91.5	9.6	641	9	AU127824	AU127824 AU127824
C 43	91.5	9.6	691	10	BB632604	BB632604 BB632604
C 44	91.5	9.6	701	14	CD262790	CD262790 PSMO19XF
C 45	91.5	9.6	844	12	B1198486	B1198486 602760491

ALIGNMENTS

RESULT 1
BF304699/c
LOCUS 601888252F1 NH_MGC_17 Homo sapiens cDNA clone IMAGE:4122276 5',
DEFINITION 884 bp mRNA linear EST 21-NOV-2000
ACCESSION BF304699.1 GI:11251586
VERSION BF304699.1
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 984)


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Db      751 AAAAAACCCAGCTGGCGCCACCGAGGCGCTAGCCCTTACAGCCAGGCGCGGG 810
      |||
QY      141 yGlyProLeuLeuCysProAlaGlyHisAlaValGlyIlePheArgAlaAlaValCysTh 162
      |||
Db      811 CCCCCCCTAACATCTCCATCCCTACCCCTGGCGCGCGGGGAGAGCTGGGCGCATACGGGC 870
      |||
QY      162 rArgGlyValaLsAlaValAlaAspPheIleProValGluSer 175
      |||
Db      871 TCAGGCGCTTTTAAAGCCCGGCGCTTCGCGCGCGCGCGAAGCA 913
      |||

RESULT 3
CA023748
LOCUS   CA023748
DEFINITION HZ47E17: HZ Hordeum vulgare subsp. vulgare cDNA clone HZ47E17
5-PRIME mRNA sequence.
ACCESSION CA023748
VERSION   CA023748.1 GI:24301122
KEYWORDS EST.
SOURCE    Hordeum vulgare subsp. vulgare
ORGANISM  Hordeum vulgare subsp. vulgare
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Poaceae
; Triticeae; Hordeum.
REFERENCE 1 (bases 1 to 515)
AUTHORS   Radchuk,V., Zhang,H., Weschke,W., Potokina,E. and Wobus,U.
TITLE     Barley ESTs from developing seeds
JOURNAL   Unpublished
COMMENT   Contact: Steig Nils
Molecular Markers Group, Department Genbank
Institute of Plant Genetics and Crop Plant Research (IPK)
Corrensstr. 3, 06466, Gatersleben, Germany
Tel: 039482-5522
Fax: 039482-5595
Email: steinipk@gatersleben.de
Insert Length: 515 Std Error: 0.00
Plate: 47 row: E column: 17
Seq primer: M3rev.
FEATURES             Location/Qualifiers
     source            1..515
     organism="Hordeum vulgare subsp. vulgare"
     mol_type="mRNA"
     cultivar="barke"
     db_xref="GABI:271158"
     /db_xref="taxon:112509"
     /clone="HZ47E17"
     /misc_type="pericarp"
     /dev_stage="0-7 DAP (days after pollination)"
     /lab_host="XL10-Gold"
     /clone_lib="H2"
     /note="Vector: pBluescript SK(+); Site1: EcoRI (5'-end of
cDNA); Site2: XhoI (3'-end of cDNA); pericarp 3-7
DAP(days after pollination). Due to a cloning artifact
caused by the kit, in most cases the EcoRI site is NOT
present, as well as the EcoRI adapter used for cloning. To
excise the insert, restriction sites upstream EcoRI should
be used (e.g. BamHI, SalI, PstI). NOTE: Also due to the
cloning system used Blue/White selection for recombinats
is not 100% reliable.Average insert size is 900 bp"
BASE COUNT      82 a 181 c 150 g 92 t
ORIGIN
Alignment scores:
Pred. No.:      5/6      Length:      515
Score:          102.00   Matches:      44
Percent Similarity: 41.96% Conservative: 16
Best Local Similarity: 30.77% Mismatches: 53
Query Match:     10.60% Gaps:      3
DB:              14      Gaps:      6

US-09-965-594-1 (1-182) x CA023748 (1-515)
QY      8 AlaGlnGlnThrArgGlyLeuLeuGlyCysIleIleThrSerLeuThrGlyArgAspLys 27

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Db      70 GCACAGCGCGCGCGCGCTAGCTAGCTGATTCGACTGTCCACCACCCAGCCACCAI 129
      |||
QY      28 AsnGlnValGlu-----GlyGluValGlnIleValSer 38
      |||
Db      130 CATCAATATCC-TCCGGCGAGAGTTTCAGCGATATTTCTGGAGAGCTTCGCGGAAGGA 188
      |||
QY      39 ThrAlaAlaGlnThrPheLeuAlaThrCysIleAsnGlyValCysTrpThrVal----- 56
      |||
Db      189 GGAGCGGCG-----GCATGGTGITCTTCTGCTCTCTCTCTCTCTCTCTCTCTCTCTCA 236
      |||
QY      57 TyrHisGlyAlaGlyThrArgTrpIleAlaSerProLysGlyProValIleGlnMetTyr 76
      |||
Db      237 AGGTGGCGGAGCAGCAGCAGCGCGGCGCATCTGCTCGCGGTGCGGGCGGTGCGCCAGCG 296
      |||
QY      77 ThrAsnValAspLysAspLeuValGlyTrpProAlaProGlnGlySerArgSerLeuThr 96
      |||
Db      297 TGGCGGACA-----TGGAGAGCGGCCACCGGCTCTGTACCTGCTGTACCTGCTGCA 338
      |||
QY      97 ProCysThrCys-----GlySerSerAspLeuValThrArgHisAlaAspVal 114
      |||
Db      339 CCGTGCACCGCGCTACCTGCGCGGCCCATCATCTGCACCTCTCTGCGGCGCATGCTCAACT 398
      |||
QY      114 IleProValArgArgArgGlyAspSerArgGlySerLeuLeuSerProArgProIleSe 134
      |||
Db      399 CCAACCG-----CCACTACCGGCTCTACTAGCCGCGGCGGTGCTGTGAC 440
      |||
QY      134 rTyrIle 136
      |||
Db      441 GCTGATT 447

BM374064      533 bp      mRNA      linear      EST 23-JUL-2002
EBma03_SQ003_I16_R maternal, 8 DPA, no treatment, cv Optic, EBma03
Hordeum vulgare subsp. vulgare cDNA clone EBma03_SQ003_I16 5', mRNA
sequence.
ACCESSION     BM374064
VERSION       BM374064.2 GI:21941288
KEYWORDS      EST.
SOURCE        Hordeum vulgare subsp. vulgare
ORGANISM      Hordeum vulgare subsp. vulgare
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Poaceae
; Triticeae; Hordeum.
REFERENCE     1 (bases 1 to 533)
AUTHORS       Hedley,P., Liu,H., Caldwell,D., McCallum,N., Mudie,S., Cardie,I.,
Ramsay,L., Machray,G., Marshall,D.F.M. and Waugh,R.
TITLE         Development of Barley Transcriptome Resources
JOURNAL       Unpublished
COMMENT       On Jan 10, 2002 this sequence version replaced gi:18117454.
Contact: Waugh R, Marshall DF
Genome Dynamics/Computational Biology
Scottish Crop Research Institute
Invergowrie, Dundee, DD2 5DA, Scotland, UK
Tel: 00 44 1382 562731
Fax: 00 44 1382 562426
Email: est@scri.sari.ac.uk
All sequence has a Phred quality score of 20 or over
Seq primer: M13 reverse.
FEATURES             Location/Qualifiers
     source            1..583
     organism="Hordeum vulgare subsp. vulgare"
     mol_type="mRNA"
     cultivar="Optic"
     db_xref="taxon:112509"
     /clone="EBma03_SQ003_I16"
     /tissue_type="maternal"
     /dev_stage="8 DPA"
     /lab_host="DH10B"
     /clone_lib="maternal, 8 DPA, no treatment, cv Optic,
EBma03"
     /note="Vector: pSPORT1; Site_1: Sal I; Site_2: Not I;

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FEATURES

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organism="Crinia laevis"
mol_type="nrna"
strain="HD-18"
db_xref="taxon:8990"
close="MP0ISSA025C02"
sex="mixture of female and
tissue_type="whole embryo"
dev_stage="segmentation stage"
clone_id="MP0ISSA CDNA"
140.0 166.0 144.0
BASE COUNT

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/clone="MP01SSA05C02"
/sex="mixture of female and male"
/tissue_type="whole embryo"
/dev_stage="segmentation stage 20"
/clone_lib="MP01SSA cDNA"
140 a 166 c 165 g 144"
BASE COUNT

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Alignment Scores:
Pred. No.:      11      Length:      615
Score:          99.00    Matches:      42
Percent Similarity: 33.77% Conservative: 9
Best Local Similarity: 27.81% Mismatches:    50
Query Match:      10.39% Indels:        50
DS:              12     Gaps:         7

US-09-965-594-1 (1-182) x B4001625 (1-615)

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Pred. NO.:	Length:	613
Score:	Matches:	42
Percent Similarity:	Conservative:	9
Best Local Similarity:	Mismatches:	50
Query Match:	Indels:	50
DB:	Gaps:	7
US-09-965-594-1 (1-182)	x	B4001625 (1-615)

U.S.-09-965-594-1 (1-182) x BJ001625 (1-615)

27 LysAsnGlnValGluC'ylGluValGlnIleValSerThrAlaAlaGlnThrPheLeuAla 46

DB 511 AAAAATGACGTAGAACCAAAAGACACAGATCCAAACCACACATGTTCTGGTGTCTACGGGCT 452

47 ThrCysIleAsnGlyValCysTrpThrValTyrHisGlyAlaGlyThrArgThrIleAla 66

Db 451 -----TGTGGAGAACCTATCACAGTTCCTGCTTTAGAGCAACGGCA 410

67 SerProLys-----GlyProValIleGlnMetTyrThrAsnValAspLys 81

DB 409 GCTCTGGCGGGCGGAGGAGCTCTGGCCACTGTG----- 374

QY 82 AspleuValGlyTrpProAlaProGlnGlySerArgSerLeuThrPro----- 97

D₁₀ 373 -----ACTCGTGGAGGACGAAGAGCGTACCCCGGAGCTGTAGG 3155

98 -----CysThrCysGlySerSerAspLeuTyrLeuValThrArg----- 110
Qy

Db
334 CTGCAGGATCGCGGATGTGGCTCTGTCT-----TTGGTTCCTCGTCTCCIGGATCA 284

Cy 111 -----HisAlaAspValIleProValArgArgArgGlyAspSer 123

25b 263 TCTTCTCAGCTGACCTTCCACATCCAGGTGTCGCCAGCGCTGGTCTGACGGGTGATGG 224

QY 124 ArgGlySerLeuLeuSerProArg-----ProIleSerTyrLeuLysGlySerSer 140

223 AGAGCGCGACAGGACGAGTCGGGGTGAATCTCTGCAGGACGTCITCAGCGGGATCA 164

QY
141 GlyCysProLeuLeuCysProAlaGlyHisAla 151

163 GGAGGACCGACTCGGTGCAGAGCCTCTGCTGCA 131

RESULT 6
EXTENDING

LOCUS	BJ024121	643 bp	mRNA	linear	EST 05-DEC-2000
DEFINITION	BJ024121	MS:SSA cDNA	Genus: latinos	cDNA	clone M805581.2012.24

DEFINITION
EU024121 M02336 CDNA Oryzias latipes CDNA M02336143D12 3
miRNA sequence.
ACCESSION
EU024121

REVISIONS	EST
VERSION	RJ024121.1
ACCESSION	BJ024121
GI	17377389

KEYWORDS: EST, *Oryzias latipes* (Japanese medaka)

Craniom
 Myzias rapides
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Actinopterygii; Neopterygii; Melosteoi; Euteleostei;

Acanthomorpha; Acanthopterygii; Percomorpha; Atherinomorpha;
Acinopterygii, Neopterygii; Teleostei; Euteleostei; Neoteleostei;
Poliopterygii, Adactynopterygii, Oxyuroidei, Gymnopteri.

Belontiiformes; Atherinomorpha; Cyprinidae; Cyprinidae; Cyprinidae.

```

REFERENCE 1 (bases 1 to 643)
AUTHORS Kohara,Y., Shin-I,T., Kimura,T., Narita,T., Jindo,T. and Takeda,H.
TITLE Medaka EST Project in Takeda's lab
JOURNAL Unpublished
COMMENT Contact: Tadasu Shin-I
Center For Genetic Resource Information
National Institute of Genetics
1111 Yata, Mishima, Shizuoka 411-8540, Japan
Tel: 81-559-81-6856
Fax: 81-559-81-6855
Email: tshini@genes.nig.ac.jp.

FEATURES
    source
        1..643
            /organism="Oryzias latipes"
            /mol_type="mRNA"
            /strain="Hd-rp"
            /db_xref="taxon:8090"
            /clone="MF01SSA143012"
            /sex="mixture of female and male"
            /tissue_type="whole embryo"
            /dev_stage="segmentation stage 20 - 25"
            /clone_lib="MF01SSA cDNA"
            /catalytic="171 a 148 c 148 g 176 t"

BASE COUNT 171 a 148 c 148 g 176 t
ORIGIN
Alignment Scores:
Pred. No.: 11.7 Length: 643
Score: 99.00 Matches: 42
Percent Similarity: 33.77% Conservatives: 9
Best Local Similarity: 27.81% Mismatches: 50
Query Match: 10.39% Indels: 50
DB: 12 Gaps: 7

US-09-965-594-1 (1-182) x BJ016176 (1-643)
QY 27 TysAsnGlnValGluGlyGluValGlnIleValSerThrAlaAlaGlnThrPheLeuAla 46
    ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 242 AAAAATGAGTAAACCAACCAACCAACCAACCAACCAACCAACCAACCAACCAACCA 302
QY 47 ThrCysIleAsnGlyValCysTrpThrValTyrHisGlyAlaGlyThrAlaThrIleAla 66
    ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 302 -----TCTGGAGAACCTATACAGTTCCTCTTACAGCAACGCCA 343
QY 67 SerProLys-----GlyProValIleGlnMetTyrThrAsnValAspLys 81
    ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 344 GCTCTCGCGCGGAGGAGTCTCTGGCCAGCTGTTG-----TCTGCTCTGCTCTGCTGCT 379
QY 82 AspLeuValGlyTrpProAlaProGlnGlySerArgSerLeuThrPro----- 97
    ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 380 -----ACTCGTGGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 418
QY 98 -----CysThrCysGlySerSerAspLeuTyrLeuValThrArg----- 110
    ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 419 CTCACGGATGCGGATGCGGCTCTCT-----TTGGTTCCTGCTCTCTCTCTCTCTCT 469
QY 111 -----HisAlaAspValIleProValAlaArgArgGlyAspSer 123
    ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 470 TCTTCTCACCCTGACCTCCACATCCAGGTGTCGCCAGCCCTGGTCTGACGGGTGATGG 529
QY 124 ArgGlySerLeuLeuSerProArg-----ProIleSerTyrLeuLysGlySerSer 140
    ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 530 AGAGCCCGACAGCGGAGTCTGGGGTGAATCTCTGAGGACCTCTTACGGCGGATCA 589
QY 141 GlyGlyProLeuLeuCysProAlaGlyHisAla 151
    ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 590 GGAGGACCGACTCGCTGCACAGCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCT 622

RESULT 7
LOCUS BJ016176 754 bp mRNA linear EST 05-DEC-2001
DEFINITION BJ016176 MF01SSA cDNA Oryzias latipes cDNA clone MF01SSA025C02 3',
mRNA sequence.
ACCESSION BJ016176

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Db      452 T-G-----GGCACACTGTGGTCCGTGGCACAT-----ATCGATGGCCCT 490
QY      69 LysGlyProValIleGlnMetIyrThrAsnValAspLysAspLeuValGlyTyrProAla 88
      ||||| ||| : : : : : : : : : : : : : : : : : : : : : : : :
Db      491 AAAGGCGCTTTCACAAAACACTT-----ACCTCCTTGGTGGCCCTGGC 535
QY      89 Pro-----GlnGlySerArgSerLeuThrProCysThrCysGly 101
      ||||| ||| : : : : : : : : : : : : : : : : : : : : : : : :
Db      536 A-T-GTTGGCCAAAAGAACCGCTTTTGGCTTGGCGCTTGGCGCCGCCCAATTGGGA 594
QY      102 SerSerAspLeuTyrLeuValThrArgHisAlaAsp-ValIleProValArgArgGly 121
      ||||| ||| : : : : : : : : : : : : : : : : : : : : : : : :
Db      595 ACCAGTGGC-----ACCACCATGGGCTTTGTGTGTCCTCCCTCCCGTGG 642
QY      121 YAspSerArgGlySerLeuLeuSerProArgProIleCysTyrLeuLysGlySerGln 141
      ||| : : : : : : : : : : : : : : : : : : : : : : : :
Db      643 GCAATTACAAACNCCCTTAACCGTCCCTCGACACAAATTTCTTANGGCTCTCTGA 702
QY      141 Y-----GlyProLeuLeuLeuCysProAlaGlyHisAlaValGly 153
      ||||| ||| : : : : : : : : : : : : : : : : : : : : : : : :
Db      703 TTTCCTTAAGTCCCGCTTTTGTACCCCAACCACTTCTGACA 748
      ||||| ||| : : : : : : : : : : : : : : : : : : : : : : : :

RESULI 10
LOCUS AK060545
DEFINITION Mus musculus 7 days neonate cerebellum cDNA, RIKEN full-length
          enriched library, clone:A730082L10 product:weakly similar to zinc
          finger protein (fragment) [Mus musculus], full insert sequence.
ACCESSION AK060545
VERSION 92279253
KEYWORDS HTO; GAP trapper
SOURCE Mus musculus (house mouse)
ORGANISM Mus musculus
REFERENCE 1
AUTHORS Carninci,P. and Hayashizaki,Y.
TITLE High-efficiency full-length cDNA cloning
JOURNAL Meth. Enzymol. 303, 19-44 (1999)
MEDLINE 92279253
PUBMED 10349536
REFERENCE 2
AUTHORS Carninci,P., Shibata,K., Hayashi,N., Sugahara,Y., Shibata,K.,
          Itoh,M., Konno,H., Okazaki,Y., Muramatsu,M. and Hayashizaki,Y.
TITLE Normalization and sub-traction of cap-trapper-selected cDNAs to
          prepare full-length cDNA libraries for rapid discovery of new genes
JOURNAL Genome Res. 10 (10), 1517-1530 (2000)
MEDLINE 20499174
PUBMED 11042159
REFERENCE 3
AUTHORS Shibata,K., Itoh,M., Aizawa,K., Nagaoka,S., Sasaki,N., Carninci,P.,
          Konno,H., Akiyama,J., Nishi,K., Kitsuami,T., Tashiro,H., Itoh,M.,
          Sumi,N., Ishii,Y., Nakamura,S., Hazama,K., Nishino,I., Harada,A.,
          Yamamoto,K., Matsumoto,H., Sakaguchi,S., Ikegami,T., Kashiwagi,K.,
          Fujiwara,S., Inoue,K., Togawa,Y., Izawa,M., Obara,S., Watahiki,M.,
          Yoneda,Y., Ishikawa,T., Ozawa,K., Tanaka,T., Matsura,S., Kawai,S.,
          Okazaki,Y., Muramatsu,M., Inoue,Y., Kira,A. and Hayashizaki,Y.
TITLE RIKEN integrated sequence analysis (RISA) system--384-format
          sequencing pipeline with 384 multicapillary sequencer
JOURNAL Genome Res. 10 (11), 1757-1771 (2000)
MEDLINE 20530913
PUBMED 11076861
REFERENCE 4
AUTHORS Kawai,J., Shibagawa,A., Shibata,K., Yoshino,M., Itoh,M., Ishii,Y.,
          Arakawa,T., Hara,A., Fukunishi,Y., Konno,H., Adachi,J., Fukuda,S.,
          Aizawa,K., Izawa,M., Nishi,K., Kiyosawa,H., Kondo,S., Yamanaka,I.,
          Saito,T., Okazaki,Y., Gojobori,T., Bono,H., Kasukawa,T., Saito,E.,
          Kadota,K., Matsuda,H., Ashburner,M., Batalov,S., Casavant,T.,
          Fleischmann,W., Gaasterland,T., Gissi,C., King,R., Kochiwa,H.,
          Kuehl,P., Lewis,S., Matsuo,Y., Nikaic,T., Pesole,G.,
          Quackenbush,J., Schraml,L.M., Staubli,F., Suzuki,K., Tomita,M.,
          Wagner,L., Washio,T., Sakai,K., Okido,T., Furuno,M., Aono,H.,
          Baldarelli,R., Barsh,G., Blake,J., Boftelli,S., Bojunga,N.,

```

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Carninci,P., de Bernaldo,M.F., Brownstein,M.J., Bult,C.,
Fletcher,C., Fujita,M., Gariboldi,M., Gustincich,S., Hill,D.,
Hofmann,M., Hume,D.A., Kamiya,M., Lee,N.H., Lyons,P.,
Marchionni,L., Mashima,J., Marzarelli,J., Mombaerts,P., Nordone,P.,
Ring,B., Ringwald,M., Rodriguez,I., Sakamoto,N., Sasaki,H.,
Sato,K., Schonbach,C., Seya,T., Shibata,Y., Storch,K.F., Suzuki,H.,
Toyooka,K., Wang,K.H., Weitz,C., Whittaker,C., Wilmang,L.,
Wynshaw-Boris,A., Yoshida,K., Hasegawa,Y., Kawaji,H., Kohsaki,S.
and Hayashizaki,Y.
Functional annotation of a full-length mouse cDNA collection
Nature 409 (6821), 685-690 (2001)
21085660
PUBMED 11217851
REFERENCE 5
AUTHORS The FANTOM Consortium and the RIKEN Genome Exploration Research
          Group Phase 1 & II Team.
TITLE Analysis of the mouse transcriptome based on functional annotation
          of 60,770 full-length cDNAs
JOURNAL Nature 420, 563-573 (2002)
6 (bases 1 to 1141)
REFERENCE Adachi,J., Aizawa,K., Akimura,T., Arakawa,T., Bono,H., Carninci,P.,
          Fukuda,S., Furuno,M., Hanagaki,T., Hara,A., Hashizume,W.,
          Hayashida,K., Hayatsu,N., Hiramoto,K., Hiraoka,T., Hirozane,I.,
          Hori,F., Imotani,K., Ishii,Y., Itoh,M., Kagawa,T., Kasukawa,T.,
          Katoh,H., Kawai,S., Kojima,Y., Kondo,S., Konno,H., Kouda,M.,
          Koya,S., Kurihara,C., Matsuyama,T., Miyazaki,A., Murata,M.,
          Nakamura,M., Nishi,K., Nomura,K., Numazaki,R., Ohno,M., Ohsato,N.,
          Okazaki,Y., Saito,K., Saitoh,H., Sakai,C., Sakai,K., Sakazume,N.,
          Sano,H., Sasaki,D., Shibata,K., Shinagawa,A., Shiraki,T.,
          Sogabe,Y., Tagami,M., Tagawa,A., Takahashi,F., Takaku-Akahira,S.,
          Takeda,Y., Tanaka,T., Tomaru,A., Toya,T., Yasunishi,A.,
          Muramatsu,M. and Hayashizaki,Y.
TITLE Direct Submission
JOURNAL Submitted (16-APR-2002) Yoshihide Hayashizaki, The Institute of
          Physical and Chemical Research (RIKEN), Laboratory for Genome
          Exploration Research Group, RIKEN Genomic Sciences Center (GSC),
          RIKEN Yokohama Institute, 1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama,
          Kanagawa 230-0045, Japan (E-mail:genome-res@gsc.riken.go.jp,
          URL:http://genome.gsc.riken.go.jp/, Tel:81-45-503-9222,
          Fax:81-45-503-9216)
COMMENT cDNA library was prepared and sequenced in Mouse Genome
          Encyclopedia Project of Genome Exploration Research Group in Riken
          Genomic Sciences Center and Genome Science Laboratory in RIKEN.
          Division of Experimental Animal Research in Riken contributed to
          prepare mouse tissues.
          Please visit our web site for further details.
          URL:http://genome.gsc.riken.go.jp/
          URL:http://fantom.gsc.riken.go.jp/
          Location/Qualifiers
            1..1141
              /organism="Mus musculus"
              /mol_type="mRNA"
              /strain="C57BL/6J"
              /db_xref="FANTOM_DB:A730082L10"
              /db_xref="taxon:10090"
              /clone="A730082L10"
              /tissue_type="cerebellum"
              /clone_lib="RIKEN full-length enriched mouse cDNA library"
              /dev_stage="7 days neonate"
            <1..587
              /note="unnamed protein product; putative
              weakly similar to zinc finger protein (fragment) [Mus
              musculus] (PIR|I48722, evidence: FASTV, 50.7%ID,
              57.6%length, match=601)"
              /codon_start=3
              /protein_id="BAC37940.1"
              /db_xref="GI:26348601"
              /translation="DSCLPAAAGRSLLTPRGDGFLEKLSAARAVGPGSVAFGV
              TVRGAQAGQRRRGACRRSGCLCLSRPRRHVPVPGVHVGLSGRRIPPPAGE
              AQAGRAPQOVPHPGPRHCTVYVPGAGLLPALAAROVPGVPGREGPRAPRHS
              PRPVPALGFSGGGPPAPLLAPANGRSVGLAL"
              1118..1123
              /note="putative"

```

polyA_signal

On Dec 19, 2000 this sequence version replaced gi:11895595.
Contact: Wing RA
Clemson University Genomics Institute
Clemson University
100 Jordan Hall, Clemson, SC 29634, USA
Tel: 864 656 7286
Fax: 864 656 4293
Email: rwing@clemson.edu
Total hg bases = 455
Seq primer: AATTAACCTCACTAAAGG
High quality sequence stop: 588.

FEATURES
source
1..779
Location/Qualifiers
/organism="Hordeum vulgare subsp. vulgare"
/mol_type="mRNA"
/cultivar="Morex"
/db_xref="taxon:112509"
/clone="HVSMB0015P05f"
/issue_type="Seedling shoot"
/lab_host="TJC121"
/clone_lib="Hordeum vulgare seedling shoot EST library
HVCDA0002 (Dehydration stress)"
/note="Vector: lambdaZAP; Site_1: EcoRI; Site_2: XhoI;
Seeds were surface sterilized then germinated under axenic
conditions in the dark at room temperature on filter paper
with water, nystatin and cefotaxime in covered
crystallization dishes. Five-day old seedlings were
incubated at 90% RH for 24 hr. Shoots were then harvested,
total RNA was prepared, poly(A) RNA was purified, one
primary unamplified cDNA library was made, 600000 pfu were
in vivo excised to give pBluescript SK(-) cDNA phagemids.
These steps were performed in the TJ Close laboratory at
the University of California, Riverside (Choi, Close,
Penton). Phagemids were plated and picked at the Clemson
University Genomics Institute (CUGI) (Begum, Palmer,
Frisch, Atkins and Wing). Plasmid DNA preparations, DNA
sequencing and sequence analysis were performed at CUGI
(Wing, Yu, Frisch, Henry, Simmons, Oates, Rambo, Main).
The sequence has been trimmed to remove vector sequence
and contains a minimum of 100 bases of phred value 20 or
above. For more details on library preparation and
sequence analysis see
http://www.genome.clemson.edu/projects/barley. To order
this clone see http://www.genome.clemson.edu/orders Also
see Close TJ, Wing R, Kleinhofs A, Wise R (2001)
Genetically and physically anchored EST resources for
barley genomics. Barley Genetics Newsletter 31:29-30.
(http://wheat.pw.usda.gov/ggpages/bgn/31/cover.html)"

BASE COUNT 147 a 216 c 311 g 105 t
ORIGIN

Alignment Scores:
Prod. No.: 21 Length: 779
Score: 97.50 Matches: 47
Percent Similarity: 39.87% Conservative: 14
Best Local Similarity: 30.72% Mismatches: 63
Query Match: 10.24% Indels: 30
DB: 10 Gaps: 6

US-09-965-594-1 (1-182) x BF631437 (1-779)

QY 35 GlnIleValSerThrAlaAlaGlnThrPheLeuAlaThrCysIleAsnGlyValCysTrp 54
||||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :||

```

QY 88 AlaProGlnGlySerArgSer-----LeuThrProCysThrCysCysSerSer 103
Db 169 TCGGGGGGTGGACCGGAGTCTGGGGCGGACGACGCTGCTGACGTGTACGACTTC 228
QY 104 AspLeuTyrLeuValThrArg-----HisAlaAspValIleProVal-----ArgAr 119
Db 229 TGACAGGCGGTGGCGGAGCGAGGCGGCCGACGTCGCGGCGGCGCGCTCTTCGCCA 288
QY 119 GArGlyAspSerArgSylSerLeuLeuSerProArgProIleSerTyrLeuLeuGlySe 139
Db 289 CCGGGCGGTTCGGGGGCGAGTATAGC-TCGCGGGCGGCGACGACGACGACGACGAG 347
QY 139 rSerGly-GlyProLeuLeuCysProAla-----GlyHisA 151
Db 348 CTCGGGTCCGGCTCGCGTACGACGACGCGGACTCCGACGGGTGGCGGCGCTCCGCGA 407
QY 151 lavalGlyIlePheArgAlaAlaValCysThrArg 162
Db 408 GCGGAGGAGCGGACGACGACGCGCGGCGGCTCGCT 442

RESULT 12
CNS03D4S/C
LOCUS
DEFINITION
Single read from an extremity of a full-length cDNA clone made from
Anopheles gambiae total adult females, 3-PRIME end of clone
FKJAC30C12 of strain 6-9 of Anopheles gambiae (African malaria
mosquito).
ACCESSION
BX053096
VERSION
BX053096.1 GI:27426377
KEYWORDS
HTC.
ORGANISM
Anopheles gambiae (African malaria mosquito)
Eukaryota; Metazoa; Hexapoda; Insecta; Pterygota;
Neoptera; Endopterygota; Diptera; Nematocera; Culicoidae;
Anopheles.
REFERENCE
1 (bases 1 to 844)
Genoscope.
Direct Submission
Submitted (05-JAN-2003) Genoscope - Centre National de Sequençage :
BP 191 91006 EVRY cedex - FRANCE (E-mail : seqref@genoscope.cns.fr)
JOURNAL
- Web : www.genoscope.cns.fr
FEATURES
Source
Location/Qualifiers
i 844
/organism="Anopheles gambiae"
/mol_type="mRNA"
/strain="6-9"
/db_xref="taxon:7145"
/clone="FKJAC30C12"
/plasmid="pXELHS-FL"
/notes="end : 3-PRIME"
BASE COUNT 197 a 220 c 213 g 214 t
ORIGIN

Alignment Scores:
Pred. No.: 24 Length: 844
Score: 96.50 Matches: 41
Percent Similarity: 35.87% Conservative: 23
Best Local Similarity: 23.37% Mismatches: 63
Query Match: 10.13% Indels: 55
DB: 11 Gaps: 10

US-09-965-594-1 (1-182) x CNS05D4S (1-844)

QY 10 GlnThrArgGlyLeuLeuGlyCysIleIleThrSerLeuThrGlyArgAspHisAsnGln 29
Db 743 CGCAGGAAGTATAACAAGCTCGTAAATTCATCTCTCCGCAACCTGGCTCGCAGAG 684
QY 30 ValGluGly-----GluValGlnIleValSerThrAlaAlaGlnThrPheLeuAlaThr 47
Db 683 CTGGCGGCGTGGGACGACGACGACGACGACGACGACGACGACGACGACGACGACGAC 624
QY 48 CysIleAsnGlyValCysTrpThrValTyrHisGlyAla-----GlyThrArgThrIle 65

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Db 623 ATCTTACCGGTGGCTGTGGCACTTCTTACAGCTGGCTGGATCGATCATCAGACGATA 564
QY 66 AlaSerProLys-----GlyProValIleGlnMetIyrThr 77
Db 563 CCCGTTCAAGAACTCTCTATGCTGCTGGCGGAGCGGCGCTTGTGGC 513
QY 78 AsnValAspLysAspLeuValGlyTrpProAlaProGlnGlySerArgSerLeuThrPro 97
Db 512 -----CGGTCAITACCGGCC 498
QY 98 CysThrCys-----GlySerSerAspLeuTyrLeuValThrArgHisAlaAsp 113
Db 497 GTGCTGTACTTGTGTCTCCGCTCGAAT-----ACAAACACACAG 456
QY 114 ValIleProValArgArgGlyAspSerArgGlySerLeuLeuSerProArgThrIle 133
Db 455 -----ACCAAGCGTCCGAATCTGTTCCACCCAGTATCGTA 420
QY 134 SerTyrLeuLysGlySerSerGlyGlyProLeuLeuCysProAlaGlyHisAlaValGly 153
Db 419 GCATTCTGC-----GGGCAACTGGAAGTCCCTTACCCGCCCTTATTCATCAACTCAACT 363
QY 154 ILePheArgAlaAlaValCysThrArgGlyValAlaLys-----AlaValAsp 169
Db 362 ACATTCCGCGA-----TGTACGCGTGTGTTGCCACATCATCGGATTCGTGGAT 309
QY 170 PheIleProVal 173
Db 308 GGTGTCTCTGTC 297

RESULT 13
BF863244
LOCUS
DEFINITION
701 bp mRNA linear EST 19-JAN-2001
963042602.xl C. reinhardtii CC-1690, Stress condition 2, normalized
Lambda Zap II Chlamydomonas reinhardtii cDNA, mRNA sequence.
ACCESSION
BF863244
VERSION
BF863244.1 GI:12253388
KEYWORDS
EST.
SOURCE
Chlamydomonas reinhardtii
ORGANISM
Chlamydomonas reinhardtii
Eukaryota; Viridiplantae; Chlorophyta; Chlorophyceae; Volvocales;
Chlamydomonadales; Chlamydomonas.
REFERENCE
1 (bases 1 to 701)
Grossman, A., Davies, J., Federspiel, N., Harris, E., Hauser, C.,
Lefebvre, P., McQuinn, J. P., Shrager, J., Sillflow, C. and Stern, D.
Analyses of the Chlamydomonas reinhardtii Genome: A Model
Unicellular System for Analyzing Gene Function and Regulation in
Vascular Plants; project phase 3
Unpublished
Contact: Charles Hauser
DCMB Box 91000
Duke University
Durham, NC 27708-1000
Tel: 919 613 8159
Fax: 919 613 8177
Email: chauser@duke.edu.
FEATURES
Location/Qualifiers
1..701
/organism="Chlamydomonas reinhardtii"
/mol_type="mRNA"
/strain="CC-1690 wild type mt+ 21gr"
/db_xref="taxon:3055"
/note="Vector: pBluescript II SK-; Site 1: EcoRI; Site 2:
XhoI; This library, constructed by John Davies and Jeffrey
McQuinn, combines cDNAs from CC-1690 cells grown to
mid-log phase in TAP-N (30 min, 1hr, 4hr), TAP-S (30 min,
1hr, 4hr), TAP-P (4hr, 12hr, 24hr), NO3 to NH4 (30min, 1hr
4hr) and NH4 to NO3 (30min, 1hr, 4hr). PolyA mRNA was
purified from each sample, pooled and cDNA synthesized.
The cDNA was directionally cloned into lambda Zap II
(Stratagene) in the EcoRI (5') and XhoI (3') sites.

```


DNA Sequencing by: Incyte Genomics, Inc.
 Clone distribution: MGC clone distribution information can be
 found through the I.M.A.G.E. Consortium/LLNL at: image.llnl.gov
 Plate: LICM1044 row: c column: 02
 High quality sequence start: 6
 High quality sequence stop: 684.

FEATURES

Location/Qualifiers
 1..901
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 /db_xref="taxon:9606"
 /clone="IMAGE:4137145"
 /tissue_type="rhabdomyosarcoma"
 /lab_host="p810B (phage-resistant)"
 /clone_lib="NIH_MGC_13"

/note="Organ: muscle; Vector: pOTB7; Site: 1: EcoRI;
 Site: 2: XhoI; cDNA made by oligo-dT priming.
 Directionally cloned into EcoRI/XhoI sites using the
 following 5' adaptor: GGCACGAG(G). Size-selected >500bp
 for average insert size 1.8kb. Library constructed by
 Ling Hong in the laboratory of Gerald M. Rubin (University
 of California, Berkeley) using ZAP-cDNA synthesis kit
 (Stratagene) and Superscript II RT (Life Technologies)."

BASE COUNT 144 a 267 c 329 q 161 t

ORIGIN

Alignment Scores:
 Pred. No.: 39.5 Length: 901
 Score: 95.50 Matches: 37
 Percent Similarity: 38.46% Conservative: 8
 Best Local Similarity: 31.62% Mismatches: 28
 Query Match: 10.02% Indels: 45
 DB: 10 Gaps: 5

US-09-965-594-1 (1-182) x BF307233 (1-901)

QY 52 ValCysTrpIhrValIyrHis-----GlyAlaGlyThrArgThrIleAl 66
 Db 620 ATGTGCTGGACGGGTCCCGCCGCGCATCTCAGCGGGGGTCCGGCAGCCACAGCGTGG 679
 QY 66 aSerProLysGlyProValIleGlnMetIyrThrAsnValAspLysAspLeuValGlyTr 86
 Db 680 TGGGCAGGACGGTGGAGTGTGCA-----GTCACAGGATG 715
 QY 86 pProAlaProGlnGlySerArgSerLeuThrProCysThrCysGlySerSerAspLeuTy 106
 Db 716 GCCCGGCCCATCCGGG----- 731
 QY 106 rLeuValIhrArgHisAlaAspValIleProValArgAlaGlyAspSerArgGlySe 126
 Db 732 -----GTGAGCCTTCCTCICAGGGGGTTCGGCGGGGGTTC 766
 QY 126 rLeuLeuSerProArgProIleSerTyrIleLysGlySerSerGlyGlyProIleLeuCy 146
 Db 767 CATTTCGGTCCCGA-----GGGGAATCTCTCCAGGGGCGCTGGACTG 810
 QY 146 sProAlaGlyHisAlaValGly-----IlePheArgAlaAlaValCys 160
 Db 811 TCCGGCGGGTTCGGCGGGCGCCACAGCGGTCCGGCGGCTGTGC 859

Search completed: September 27, 2003, 14:09:39
 Job time : 2238 secs

SUMMARY

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed.

```

/*tag- a
/product= "HCV-1 NS3/4a conformational antigen"
/Note= "CDS does not include stop codon"
/partial

WO200196875-A2.
20-DEC-2001.

14-JUN-2001: 2001WO-US19169.
15-JUN-2000: 2000US-212082P.
02-APR-2001: 2001US-280811P.
02-APR-2001: 2001US-280867P.
(CHIR ) CHIRON CORP.

Chien DY, Arcangel P, Tandeske L, George-Nascimento C, Coit D;
Medina-Selby A;
WPI: 2002-179522/23.
P-PSDB; AAE18689.

Immunosay solid support useful for detecting hepatitis C virus
infection in a biological sample, comprises at least one of HCV
anti-core antibody and HCV NS3/4a epitope, bound to the support .

Example 2: Fig 4; 37pp; English.

The present invention relates to hepatitis C virus (HCV) core antigen
and NS (nonstructural) 3/4a antibody combination assay that can detect
both HCV antigens and antibodies present in a sample using a single
solid matrix as well as immunosay solid supports for use in the assay.
The solid support is useful for detecting HCV infection in a biological
sample. The present sequence is a DNA encoding HCV-1 NS3/4a mutant
conformational antigen. This sequence is used in the exemplification
of the invention.

Sequence 2058 BP: 419 A: 634 C: 580 G: 425 T: 0 other:

Alignment Scores:
Pred. No.: 1,456-78 Length: 2058
Score: 345.00 Matches: 190
Percent Similarity: 100.00% Conservative: 2
Best Local Similarity: 98.90% Mismatches: 0
Query Match: 95.27% Indels: 0
DB: 24 Gaps: 0

us-09-965-594-1 (1-182) x AAD29795 (1-2058)

QY 1 MetAlaProIleThrAlaTyAlaGlnGlnThrArgGlyLeuLeuGlyCysIleIleThr 20
DB 1 ATFGCGCCCATCACGGCGTAGCCCGAGAGCAAGGGGCCCTCTAGGGTGCATATCACC 60
QY 21 SerLeuThrGlyArgAspGlyAsnGlnValGlnGlyGluValGlnIleValSerThrAla 40
DB 61 AGCCTAACTGGCGCGGCAAAACCAAGTGGAGGTGAGTCCAGATTGTCTCAACTGCT 120
QY 41 AlaGlnThrPheLeuAlaThrCysIleAsnGlyValCysTrpIleValIleHisLeuAla 60
DB 121 GCCCAAACTTCTGGCAACGTGCAATCAATGGGTGTGTGACGTGTATCCACGGGGCC 180
QY 61 GlyThrArgThrIleAlaSerProLysGlyProValIleGlnMetTyThrAsnValAsp 80
DB 181 GGACGAGGACCATCGCTCACCCAAAGGGTCTGTGTATCCAGATGTATACCAATGTAGAC 240
QY 81 LysAspLeuValGlyTrpProAlaProGlnGlySerArgSerLeuThrProCysThrCys 100
DB 241 CAAGACCTGTGGGTGGCGCCCTCCGCAAGTACCGCATTCATTCACACCTGCACATCC 300
QY 101 GlySerSerAspLeuTyIleuValThrArgHHisAlaAspValIleProValArgArgArg 120
DB 301 GGCTCTCGGACCTTTACCTGTGTCAGGAGGACGCCGATGTCTTCCGTGCGGGCGG 360

QY 121 GlyAspSerArgGlySerLeuLeuSerProArgProIleSerTyLeuLeuYSGYSerSer 140
DB 361 GGTGATAGCAGGGGCGAGCGTGTGCGCCCGGCCCATTTCTTACITGAAGGCTCTCIG 420
QY 141 GlyGlyProLeuLeuCysProAlaGlyHisAlaValIleGlyPheArgAlaAlaValCys 160
DB 421 GGGGCTCCGCTGTGTGCGCCCGGGGCGACCGCGTGGCATATTTAGGGCGCGGTGTGC 480
QY 161 ThrArgGlyValAlaLysAlaValAspPheIleProValGluSerLeuGluThrMet 180
DB 481 ACCCGTGGAGTGGCTAAGCGGTGGACTTATCCCTGTGGAGAACCTAGAGACACCATG 540
QY 181 ArgSer 182
DB 541 AGGTCC 546

RESULT 2
ABK15344
ID ABK15344 standard; DNA: 2058 BP.
XX
AC ABK15344.
XX
DE 08-MAY-2002 (first entry)
DE Hepatitis C virus NS3/4a conformational epitope gene sequence.
XX
KW Hepatitis C virus; HCV; NS3/4a conformational epitope; seroconversion;
KW immunosay solid support; multiple epitope fusion antigen; MEFA;
KW non-structural protein; gene; ds.
XX
OS Hepatitis C virus.
XX
FH Key Location/Qualifiers
FT CDS 1..2058
FT /tag- a
FT /product= "HCV NS3/4a conformational epitope"
FT /note= "This sequence lacks a stop codon"
XX
XX WO200196870-A2.
XX
XX 20-DEC-2001.
XX
XX 14-JUN-2001; 2001WO-US19156.
XX
XX 15-JUN-2000; 2000US-212082P.
XX
XX 02-APR-2001; 2001US-280811P.
XX
XX 02-APR-2001; 2001US-280867P.
XX
XX (CHIR ) CHIRON CORP.
XX
XX Chien DY, Arcangel P, Tandeske L, George-Nascimento C, Coit D;
XX Medina-Selby A;
XX
XX WPI: 2002-090228/12.
XX
XX P-PSDB; AAU76377.
XX
XX Immunosay solid support, useful for detecting hepatitis C virus
XX infection in biological sample, comprises HCV NS3/4a conformational
XX epitope and multiple epitope fusion antigen bound to the support -
XX
XX Disclosure: Fig 3; 92pp; English.
XX
XX The present invention relates to a new immunosay solid support
XX consisting essentially of at least one hepatitis C virus (HCV) NS3/4a
XX conformational epitope and a multiple epitope fusion antigen (MEFA),
XX bound to the support. The NS3/4a conformational epitope and/or
XX MEFA reacts specifically with anti-HCV antibodies present in a biological
XX sample from an HCV-infected individual. The immunoassay of the invention
XX is useful for detecting hepatitis C virus infection in a biological
XX sample. The method of the invention provides a sensitive, accurate
XX diagnostic and prognostic tool to provide adequate patient care and to

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CC prevent transmission of HCV by blood and by blood products, or by
CC personal contact. Use of NS3/4a conformational epitope in combination
CC with MEPA, provides a sensitive and reliable method for detecting early
CC HCV seroconversion. Use of MEPA has the added advantages of decreasing
CC masking problems, improving sensitivity in detecting antibodies by
CC allowing a greater number of epitopes on a unit surface area of
CC substrate, and improving substrate detection accuracy is increased and
CC the incidence of false results is reduced because of the identification
CC and the use of highly immunogenic HCV antigens which are present during
CC the early stages of HCV seroconversion. The present nucleic acid sequence
CC encodes the non-structural protein NS3/4a conformational epitope of the
CC invention.

CC
CC
CC
SQ Sequence 2058 BP: 419 A; 633 C; 561 G; 425 T; 0 other:

DE	DNA encoding HCV-1 NS3/4a conformational antigen.
XX	
XX	Immunoassay solid support; Hepatitis C Virus type-1; HCV-1;
XX	NS3/4a conformational epitope; multiple epitope fusion antigen;
KW	MHFA; anti-HCV antibody; NS3/4a conformational antigen;
KW	HCV infection; mutant; gene; ds.
XX	
OS	Hepatitis C virus type 1.
OS	Synthetic.
EH	Key location/Qualifiers
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FT	/*tag= a
FT	/partial=
FT	/product= "NS3/4a conformational antigen"
FT	/note= "This sequence lacks a stop codon"
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PN	US2002146685-A1.
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PD	10-OCT-2002.
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PE	14-JUN-2001; 2001US-C881654.
XX	
PR	15-JUN-2000; 2000US-212082P.
PR	02-APR-2001; 2001US-280811P.
PR	02-APR-2001; 2001US-280867P.
XX	
XX	(CHIE/) CHIEN D Y.
PA	(ARCA/) ARANGEL F.
PA	(TAND/) TANDESKE L.
PA	(GEOR/) GEORGE-NASCIMENTO C.
PA	(COIT/) COIT D.
PA	(MED/) MEDINA-SELBY A.
XX	
XX	Chien DY, Arcangel P, Tandeske L, George-Nascimento C, Coit D;
PI	Medira-Selby A;
XX	
XX	WP1: 2003-147573/14.
XX	P-PSDB; ABG72261.
XX	
PI	Immunoassay solid support for detecting Hepatitis C Virus infection in
P:	biological samples, comprises Hepatitis C Virus conformational epitope
P:	and multiple epitope fusion antigen -
XX	
PS	Disclosure: Fig 3A-3D; 45pp; English.
CC	The present invention relates to immunoassays comprising Hepatitis C
CC	Virus (HCV) NS3/4a conformational epitope and multiple epitope fusion
CC	antigen (MHFA), bound to a solid support. The NS3/4a epitope and/or
CC	the multiple epitope fusion antigen react with anti-HCV antibodies
CC	present in a biological sample from an HCV-infected individual. The
CC	immunoassays and methods of the invention are useful for detecting
CC	HCV infection in a biological sample. The inventive immunoassay solid
CC	support provides a sensitive and reliable method for detecting early
CC	HCV seroconversion. The assays can detect HCV infection caused by any
CC	six known genotypes of HCV. The use of the multiple epitope fusion
CC	proteins decreases masking problems, improves sensitivity in detecting
CC	antibodies by allowing a greater number of epitopes on a unit area
CC	of substrate, and improves selectivity. The present sequence
CC	encodes HCV type 1 (HCV-1) NS3/4a conformational antigen, a mutant of
CC	the HCV-1 NS3/4a polypeptide.
XX	
SO	Sequence 2058 BP; 419 A; 633 C; 581 G; 425 T; 0 other:
Alignment Scores:	
Pred. No.:	1,466-78 Length: 2058
Score:	946.00 Matches: 180
Percent Similarity:	100.00% Conservative: 2
Best Local Similarity:	98.90% Mismatches: 0
Query Match:	99.27% Indels: 0
DB:	25 Gaps: 0
US-09-965-594-1 (1-182)x ABX14410 (1-2058)	

Qy 1 MetAlaProIleThrAlaIyrAlaGlnGlnThrArgGlyLeuLeuGlyCysIleIleThr 20
 Db ATGCGGCCCATCACGGCTACGGCCAGCAGACAGAGGGGCTCTAGGTCATATCACC 60
 Qy 21 SerLeuThrGlyArgAspLysAsnGlnValGlnGlyGluValGlnIleValSerThrAla 40
 Db AGCTTAAGTGGCGGGACAAACCAAGTGGAGGGTGGAGTCCAGATGTGTCAACTGCT 120
 Qy 41 AlaGlnThrPheLeuAlaThrCysIleAspGlyValCysIrrPthrValThrValGlyAla 60
 Db GCCCAACCTTCCTGGCAACCTGCATCAATGGGTGTCTGGACCTCTACAGGGGGCC 180
 Qy 61 GlyThrArgThrIleAlaSerProLysGlyProValIleSerMetTyrThrAsnValAsp 80
 Db GGAACGAGGACCATCGGCTCACCAAGGGTCTCTCATCCAGATGATACCAATGTAGAC 240
 Qy 81 LysAspLeuValGlyTyrProAlaProGlnGlySerArgSerLeuThrProCysThrCys 100
 Db CAGACCTTGTGGCTGGCGCGCTCCCGCAAGTAGCCGATCATTGACACCCCTGCCTTC 300
 Qy 101 GlySerSerAspLeuThrValThrArgHisAlaAspValIleProValArgArgArg 120
 Db GGCTCCCGGACCTTACCTGGTCACGAGCAGCCGCACTCATTCGCGCGCGCGG 360
 Qy 121 GlyAspSerArgGlySerLeuLeuSerProArgProIleSerTyrLeuGlySerSer 140
 Db GGTGATACGAGGAGCCCTGCTGTCCCGCGCCCATCTCTACITGAAGGCTCTCTCG 420
 Qy 141 GlyGlyProLeuLeuCysProAlaGlyHisAlaValGlyIlePheArgAlaAlaValCys 160
 Db GGGGTCGCTGTGTGGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCG 480
 Qy 161 ThrArgGlyValAlaLysAlaValAspPheIleProValGluSerLeuGluThrThrMet 180
 Db ACCCGTGGAGTGGTAGCGGTGGACCTTATCCCTGTGGAGACCTTAGAGACACCATG 540
 Qy 181 ArgSer 182
 Db AGGTCC 546

RESULT 4
 AAN92097
 ID AAN92097 standard: DNA: 5300 BP.
 AC AAN92097;
 XX
 XX 25-MAR-2003 (updated)
 DT 02-MAR-1990 (first entry)
 XX
 DE Combined open reading frames of the hepatitis C virus (HCV) cDNA in
 DE clones 141, 11b, 7f, 7e, 8h, 33c, 40b, 37b, 35, 36, 81, 32, 33b, 25c, 14c.
 DE 8f, 33f, 33g and 39c.
 KW Hepatitis C virus: HCV; non-A, non-B hepatitis; NANBH.
 XX
 OS Hepatitis C virus.
 XX
 FH Key Location/Qualifiers
 FT CDS 3..5300
 FT /tag- a
 XX
 PN EP318216-A.
 XX
 XX 31-MAY-1989.
 XX
 PF 18-NOV-1988; 88EP-0310922.
 XX
 XX 16-NOV-1987; 87US-0122714.
 PR 30-DEC-1987; 87US-0139886.
 PR 26-FEB-1988; 88US-0161072.
 PR 06-MAY-1988; 88US-0191251.
 PR 26-OCT-1988; 88US-0263584.

PR 14-NOV-1988; 88US-0271450.
 XX
 PA (CHIR) CHIRON CORP.
 XX
 PI Houghton M, Choo QL, Kuo G;
 XX
 XX MPI; 1989-159274/22.
 DR P-PSDB; AAP92041.
 XX
 XX Purified hepatitis C virus
 PI - and associated nucleic acids and polypeptide(s)
 XX
 XX Claim 3: Figure 26-1, 26-2, 26-3, 26-4, 26-5, 26-6; 139pp: English.
 XX
 XX It is a double-stranded nucleotide sequence of the open reading frame
 CC (ORF) (tag a) extending through clones 141, 11b, 7f, 7e, 8h 33c, 40b,
 CC 37b, 35, 36, 81, 32, 33b, 25c, 14c, 8f, 33f, 33g and 39c of hepatitis C
 CC virus (HCV) cDNA. In creating the composite sequence the following
 CC heterogeneities were considered. Clone 33c contains a sequence
 CC of 800 base pairs which overlaps the cDNAs in clones 40b and 37c. In
 CC clone 33c, as well as in 5 other overlapping clones, nucleotide #789 is
 CC a G. However, in clone 37b the corresponding nucleotide is an A. This
 CC heterogeneity may have important ramifications for protein folding.
 CC Nucleotide #2 in clone 8h is a T which may represent a cloning artifact
 CC because the corresponding residue in clone 7e and in 3 other overlapping
 CC clones is an A. Therefore the residue in this position is designated as
 CC an A. The 3'-terminal nucleotide in clone 8f is represented as a T
 CC than a G because the corresponding residue in clone 33f and in 2 other
 CC overlapping clones is a T. The 3' terminal sequence of clone 33f is
 CC represented as ATTC, as is found in the corresponding sequence in clone
 CC 33g and in 2 other overlapping clones, rather than as TTGC, as is found
 CC in clone 33f. Residue #4 in clone 33g is designated an A rather than a T
 CC because the corresponding residue in clone 33f and 2 other overlapping
 CC clones is an A. The 3'-terminus of clone 141 is depicted as TA rather
 CC than AA because the corresponding dinucleotide in clone 11b and 3 other
 CC clones is TA. Potential cloning artifacts have been omitted and instead
 CC the corresponding sequences in non-5'-terminal regions of multiple
 CC overlapping clones are shown. AAN92097 could be used as a source of
 CC oligomeric DNA hybridisation probes to detect the presence of HCV
 CC nucleic acids in samples. The polypeptide(s) it encodes could be used as
 CC immuno- assay reagents and vaccines and to generate antibodies useful in
 CC diagnosis and passive immunotherapy for HCV infection/non-A, non-B
 CC hepatitis.
 CC (Updated on 25-MAR-2003 to correct PR field.)
 CC (Updated on 25-MAR-2003 to correct PI field.)
 XX
 XX Sequence 5300 BP; 1047 A; 1606 C; 1515 G; 1130 T; 2 other;
 SQ
 Alignment Scores:
 Pred. No.: 8.9e-78 Length: 5300
 Score: 943.00 Matches: 179
 Percent Similarity: 100.00% Conservative: 3
 Best Local Similarity: 98.35% Mismatches: 0
 Query Match: 98.95% Indels: 0
 DB: 10 Gaps: 0
 CS-09-965-594-1 (1-162) x AAN92097 (1-5300)
 Qy 1 MetAlaProIleThrAlaIyrAlaGlnGlnThrArgGlyLeuLeuGlyCysIleIleThr 20
 Db CTGGCGCCCATCACGGCTACGGCCAGCAGACAGAGGGGCTCTAGGTCATATCACC 989
 Qy 21 SerLeuThrGlyArgAspLysAsnGlnValGlnGlyGluValGlnIleValSerThrAla 40
 Db AGCCTAACTGGCGGGCAGCAAAACCAAGTGGAGGGTGGAGTCCAGATGTGTCAACTGCT 1049
 Qy 41 AlaGlnThrPheLeuAlaThrCysIleAsnGlyValCysIrrPthrValThrValGlyAla 60
 Db GCCCAACCTTCCTGGCAACCTGCATCAATGGGTGTCTGGACCTCTACAGGGGGCC 1109
 Qy 61 GlyThrArgThrIleAlaSerProLysGlyProValIleSerMetTyrThrAsnValAsp 80
 Db GGAACGAGGACCATCGGCTCACCAAGGGTCTCTCATCCAGATGATACCAATGTAGAC 1169

QY 81 LysAspLeuValIleYrPProLafrogLinglySerArgSerLeuThrProCysThrCys 100
 ::
 Db 1170 CAAGACCTGTGGGCTGGCCGCTCGGCAAGGTAGCGGTCAATTGACACCTGCACTTGC 1229
 QY 131 GlySerSerAspLeuTyrLeuValThrArgHisAlaAspValIleProValArgArgArg 120
 ::
 Db 1230 GGCTCTCGGACCTTTACTGTGTACAGAGGACGCGGTATGTATTCCTCGCGCGGGCGG 1269
 QY 121 GlyAspSerArgGlySerLeuLeuSerProArgProLisSerTyrLeuGlyGlySerSer 140
 ::
 Db 1290 GTGTATAGCAGGGGACGCTGTGTGGCCCGGCGCAATCTGCTACTTGAAGGCTGCTGC 1349
 QY 141 GlyGlyProLeuLeuCysProAlaGlyHisAlaValGlyLeuPheArgAlaAlaValCys 160
 ::
 Db 1350 GGGGGTCCGCTGTGTGTGCGCGCGGCGGCGGCGGCGGCGGCGGCGGCGGCGGCGG 1409
 QY 161 ThrArgGlyValAlaLysAlaValAspPheIleProValGlySerLeuGluThrMet 180
 ::
 Db 1410 ACCCGTGGAGTGGCTTAGGCGGTGGACCTTATGCTGTGGAGACCTAGAGACACCATG 1469
 QY 181 ArgSer 182
 :::::
 Db 1470 AGGTCC 1475

RESULT 5
 AAN90327
 ID AAN90327 standard; cDNA; 5360 BP.
 AC AAN90327;
 DT 25-MAR-2003 (updated)
 DT 11-NOV-1989 (first entry)
 XX Hepatitis C virus composite probe.
 XX Hepatitis C virus: composite cDNA; probe; vaccine.
 XX Pan troglodytes.
 XX Key Location/Qualifiers
 FT CDS 3..5360
 FT /*tag= a
 XX GB22:2511-A.
 XX 26-JUL-1989.
 XX 18-NOV-1988; 88GB-0027024.
 XX 18-NOV-1987; 87US-0122714.
 XX 30-DEC-1987; 87US-0139886.
 XX 26-FEB-1988; 88US-0161072.
 XX 26-OCT-1988; 88US-0263584.
 XX (CHIR) CHIRON CORPORATION.
 XX Houghton M, Choo QL, Kuo G;
 XX WFI; 1989-215054/40.
 XX Hepatitis C virus gene - used for prodn. of polynucleotide probes,
 PT polypeptide(s) and antibodies for diagnosis, prevention and treatment
 PT of infection.
 PS Disclosure: Fig. 26; 174pp; English.
 CC The sequence shows the composite cDNA sequence derived from the aligned
 CC hepatitis C virus (HCV) cDNA's in clones 141, 11b, 7f, 7e, 8h, 33c, 40b,
 CC 37b, 35, 36, 81, 32, 33b, 25c, 14c, 8f, 33f, 33g and 39c. The cDNA
 CC encodes antigens which react with antibodies in patients with non-A
 CC non-B hepatitis (NANBH). The cDNA can be used to design probes, or to
 CC synthesise polypeptides, which are used to diagnose HCV-induced NANBH,

CC to raise antibodies for immunoassay or treatment, or to produce
 CC vaccines. See also AAN90158, AAN90303-26, and AAN90328-36.
 CC (Updated on 25-MAR-2003 to correct PR field.)
 XX
 SQ Sequence 5360 BP; 1060 A; 1622 C; 1532 G; 1145 T; 1 other:
 Alignment Scores:
 Pred. No.: 9, 04e-78 Length: 5360
 Score: 943.00 Matches: 179
 Percent Similarity: 100.00% Conservative: 3
 Best Local Similarity: 98.35% Mismatches: 0
 Query Match: 98.95% Indels: 0
 Gaps: 10
 DB: 10
 US-09-965-594-1 (1-182) x AAN90327 (1-5360)
 QY 1 MetAlaProIleThrAlaTyrAlaGlnGlnThrArgGlyLeuLeuGlyCysIleIleThr 20
 ::
 Db 930 CTGGGCCCATCAGGGCTAGCCAGCAGACAGAGGGGCTCTTAGGGTGCATATCACC 989
 QY 21 SerLeuThrGlyArgAspLysAsnGlnValGluGlyGluValGlnIleValSerThrAla 40
 ::
 Db 990 AGCCTAACTGGCGGGGACAAAACCAAGTGGAGGCTCAGGTCCAGANTGTGTCAACTGCT 1049
 QY 41 AlaGlnThrPheLeuAlaThrCysIleAsnGlyValCysTyrThrValTyrHisGlyAla 60
 ::
 Db 1050 GCCCAACCTTCCTGGCAACGTGCATCAATGGGTGTGCTGGACGTCTACACGGGGGCC 1109
 QY 61 GlyThrArgThrIleAlaSerProLysGlyProValIleGlnMetTyrThrAsnValAsp 80
 ::
 Db 1110 GGAAGGAGGACCATCGCGTCACCAAGGCTCTGTCTATCCAGATGTATACCAATGTAGAC 1169
 QY 81 LysAspLeuValGlyTyrProAlaProGlnGlySerArgSerLeuThrProCysThrCys 100
 ::
 Db 1170 CAAGACCTGTGGGCTGGCCGCTCGGCAAGGTAGCGGTCAATGACACCTGCACTTGC 1229
 QY 101 GlySerSerAspLeuTyrLeuValThrArgHisAlaAspValIleProValArgArgArg 120
 ::
 Db 1230 GGCTCTCGGACCTTTACTGTGTACAGAGGACGCGGTATGTATTCCTCGCGCGGGCGG 1289
 QY 121 GlyAspSerArgGlySerLeuLeuSerProArgProLisSerTyrLeuGlyGlySerSer 140
 ::
 Db 1290 GTGTATAGCAGGGGACGCTGTGTGGCCCGGCGCAATCTGCTACTTGAAGGCTGCTGC 1349
 QY 141 GlyGlyProLeuLeuCysProAlaGlyHisAlaValGlyLeuPheArgAlaAlaValCys 160
 ::
 Db 1350 GGGGGTCCGCTGTGTGTGCGCGCGGCGGCGGCGGCGGCGGCGGCGGCGGCGGCGG 1409
 QY 161 ThrArgGlyValAlaLysAlaValAspPheIleProValGlySerLeuGluThrMet 180
 ::
 Db 1410 ACCCGTGGAGTGGCTTAGGCGGTGGACCTTATGCTGTGGAGACCTAGAGACACCATG 1469
 QY 181 ArgSer 182
 :::::
 Db 1470 AGGTCC 1475

RESULT 6
 AAN92103
 ID AAN92103 standard; DNA; 6905 BP.
 XX AAN92103;
 AC AAN92103;
 XX 25-MAR-2003 (updated)
 DT 02-MAR-1990 (first entry)
 XX
 DE Combined open reading frames of the hepatitis C virus (HCV) cDNAs from
 DE clones 12f through 15e.
 XX Hepatitis C virus; HCV; non-A, non-B hepatitis; NANBH.
 XX Hepatitis C virus.
 XX Key Location/Qualifiers

```

FT CDS 3..6905
FT /*tag= a
XX EP18216-A..
XX 31-MAY-1989: 88EP-0310922.
XX 16-NOV-1988:
XX 18-NOV-1987: 87US-0122714.
XX 30-DEC-1987: 87US-0139886.
XX 26-FEB-1988: 88US-0161072.
XX 06-MAY-1988: 88US-0191263.
XX 26-OCT-1988: 88US-0263584.
XX 14-NOV-1988: 88US-0271450.
XX (CHIR ) CHIRON CORP.
XX Houghton M, Choo QL, Kuo G;
XX WPI: 1989-159274/22.
XX P-PSDB: AAP92047.
XX Purified hepatitis C virus
XX - and associated nucleic acids and polypeptide(s)
XX Claim 3; Figure 32-1 - 32-7; 139pp; English.
XX It is a double-stranded nucleotide sequence of the open reading frame
XX (ORF) (tag a) extending through clones 121 to 156 of hepatitis C virus
XX (HCV) cDNA. It can be used to make oligomeric DNA hybridisation probes to
XX detect the presence of HCV nucleic acids in samples. The polypeptide(s)
XX it encodes could be used as immunoassay reagents and vaccines and to
XX generate antibodies useful in diagnosis and passive immunotherapy for
XX HCV infection/non-A, non-B hepatitis.
XX (Updated on 25-MAR-2003 to correct PR field.)
XX (Updated on 25-MAR-2003 to correct PI field.)
XX Sequence 6905 BP; 1421 A; 2082 C; 1946 G; 1456 T; 0 other;

Alignment Scores:
Pred. No.: 1,238-77 Length: 6905
Score: 943.50 Matches: 179
Percent Similarity: 100.00% Conservative: 4
Best Local Similarity: 98.35% Mismatches: 0
Query Match: 98.95% Indels: 0
DB: 1.0 Gaps: 0

US-09-965-594-1 (1-182) x AAN92103 (1-6905)

QY 1 MetAlaProfilThrAlaTyraIaGlnGlnThrArgGlyLeuLeuGlyCysIleIleThr 20
DB 1203 CTGGCGCCATCAGGGCTACGCCAGCAGACAGAGGGGCCCTCTAGGCTGCAZANTCAGC 1262

QY 21 SerLeuThrGlyArgAspLysAsnGlnValGluGlyGlnValGlnIleValSerThraIa 40
DB 1263 AGCCTTAATGGCCGGCAAAACCAAGATGGAGGCTGAAGTCCACAGATGTGTCAACTGCT 1322

QY 41 AlaGlnThrPheLeuAlaThrCysIleAsnGlyValCysIleThrValThrHisGlyAla 60
DB 1323 GCCCAAACTCTCTGGCAACTGCTGATCAATAGGGGTGTGCTGGACTGTATACAGGGGCC 1382

QY 61 GlyThrArgThrIleAlaSerProLysGlyProValIleGlnMetIleThrAsnValAsp 80
DB 1383 GGAACGAGGACCATCGCTACCCCAAGGCTCTGTCTATCCAGATGTATACCAATGTAGAC 1442

QY 81 LysAspLeuValGlyTyrProAlaProGlnGlySerArgSerLeuThrProCysThrCys 100
DB 1443 CAAGACCTTGTGGCTGCGCCCTCCGCAAGGTATGCGCTATTGACACCCCTGCACCTGC 1502

QY 101 GlySerSerAspLeuTyrLeuValThrArgHisAlaAspValIleProValArgArgArg 120
DB 1503 GGCTCTCGACCTTACCTTGTGTCACGAGGACGCCGATGTCATTCCTCGCGCGCGGG 1562

```

```

QY 121 GlyAspSerArgGlySerLeuLeuSerProArgProIleSerTyrLeuLysGlySerSer 140
DB 1563 GGTGATAGAGGGCGCAGCCTGTGTGCGCCCGCCCATTTCTTACTTGAAGCTCCCTCG 1622

QY 141 GlyGlyProLeuLeuCysProAlaGlyHisAlaValGlyIlePheArgAlaAlaValCys 160
DB 1623 GGGGGTCCCTGTTGTGCCCCGGGCGACGGCGTGGCATATTTAGGGCCGCGGTGTC 1682

QY 161 ThrArgGlyValAlaLysAlaValAspPheIleProValGlnSerLeuGluThrThrMet 180
DB 1683 ACCCGTGGAGTGGCTAAGCGGTGGACTTATCCCTGTGAGAACCTAGAGACCAACCAT 1742

QY 181 ArgSer 182
DB 1743 AGGTCC 1748

RESULT 7
AAN92106
T3 AAN92106 standard; DNA: 7310 BP.
XX AAN92106;
XX 25-MAR-2003 (updated)
DT 02-MAR-1990 (first entry)
XX Combined open reading frames of the hepatitis C virus (HCV) cDNAs from
XX clones K9-1 through 156.
XX Hepatitis C virus: HCV; non-A, non-B hepatitis; NANBH.
XX Hepatitis C virus.
XX Key Location/Qualifiers
XX CDS 3..7310 /*tag= a
XX EP18216-A.
XX 31-MAY-1989.
XX 16-NOV-1988: 88EP-0310922.
XX 18-NOV-1987: 87US-0122714.
XX 30-DEC-1987: 87US-0139886.
XX 26-FEB-1988: 88US-0161072.
XX 06-MAY-1988: 88US-0191263.
XX 26-OCT-1988: 88US-0263584.
XX 14-NOV-1988: 88US-0271450.
XX (CHIR ) CHIRON CORP.
XX Houghton M, Choo QL, Kuo G;
XX WPI: 1989-159274/22.
XX P-PSDB: AAP92050.
XX Purified hepatitis C virus
XX - and associated nucleic acids and polypeptide(s)
XX Claim 3; Figure 47-1 - 47-8; 139pp; English.
XX It is a double-stranded nucleotide sequence of the open reading frame
XX (ORF) (tag a) extending through clones K9-1 to 156 of hepatitis C virus
XX (HCV) cDNA. It can be used to make oligomeric DNA hybridisation probes to
XX detect the presence of HCV nucleic acids in samples. The polypeptide(s)
XX it encodes could be used as immunoassay reagents and vaccines and to
XX generate antibodies useful in diagnosis and passive immunotherapy for
XX HCV infection/non-A, non-B hepatitis.
XX (Updated on 25-MAR-2003 to correct PR field.)
XX (Updated on 25-MAR-2003 to correct PI field.)
XX Sequence 7310 BP; 1491 A; 2217 C; 2058 G; 1540 T; 4 other;

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Alignment Scores:
Pred. No.: 1,32e-77 Length: 7310
Score: 943.00 Matches: 179
Percent Similarity: 100.00% Conservatives: 3
Best Local Similarity: 98.35% Mismatches: 0
Query Match: 98.95% Indels: 0
DB: 10 Gaps: 0

US-09-965-594-1 (1-182) x AAN90336 (1-7310)
QY 1 MetAlaProIleThrAlaTyrAlaGlnThrArgGlyLeuLeuGlyCysIleIleThr 20
DB 1728 CTGGCCGCCATCAGCGGTACGCCAGCAGCAGCAAGGGGCTCTAGGGTGCATATCACC 1787
QY 21 SerLeuThrGlyArgAspLysAsnGlnValGluValGlnIleValSerThrAla 40
DB 1788 AGCCTAACTGGCGGACAAAACCAAGTGGAGGGTCAGTTCAGATTTGTGCAACTGCT 1847
QY 41 AlaGlnThrPheLeuAlaThrCysIleAsnGlyValCysTrpThrValTyrHisGlyAla 60
DB 1848 GCCCAACCTTCTGGCAACGTGCATCAATGGGG:GTGGTGGACTGTCTACCCAGCGGGC 1907
QY 61 GlyThrArgThrIleAlaSerProLysGlyProValIleGlnMetTyrThrAsnValAsp 80
DB 1908 GGAACGAGGACCATCGCTCACCAAGGTCCGTGTCATCCAGATGTATACCAATGTAGAC 1967
QY 81 LysAspLeuValGlyTrpProAlaProGlnGlySerArgSerLeuThrProCysThrCys 100
DB 1968 CAAGACCTTGTGGGTGGCGCGCTCCGCAAGGTAGCGGTGTCATTCACACCTGCATTCG 2027
QY 101 GlySerSerAspLeuTyrLeuValThrArgHisAlaAspValIleProValArgAtg 120
DB 2028 GGCCTCTCGSACCTTTACCTGGTCAGGCGACGGGATGTATTCCTCCGTGGCGCGCG 2087
QY 121 GlyAspSerArgGlySerLeuLeuSerProArgProIleSerTyrLeuLysGlySerSer 140
DB 2088 GGTGATAGCAGGGCAGCCTGCTGTCCGCCGCCCATTTCTTACTTTGAAGGCTCCTCG 2147
QY 141 GlyGlyProLeuLeuCysProAlaGlyHisAlaValGlyIlePheArgAlaValCys 160
DB 2148 GGGGGTCCGCTGTGTGTGGCGGGGACCGCGGTGGCATATTTAGGGCGCGGTGTGC 2207
QY 161 ThrArgGlyValAlaLysAlaValAspPheIleProValSerLeuGlnThrMet 180
DB 2208 ACCGTGGAGTGGTAGGGCGGTGGACTTTATCTGTCTGTGAGAACTTAGACAACCATG 2267
QY 181 ArgSer 182
DB 2268 AGCTCC 2273

RESULT 8
AAN90336
ID AAN90336 standard: DNA: 7310 BP.
XX AC AAN90336;
XX AC
XX 25-MAR-2003 (updated)
DT 19-JUL-2001 (updated)
DT 01-NOV-1989 (first entry)
XX XX
DE Composite hepatitis C virus (HCV) cDNA.
KW Hepatitis C virus: cDNA: clone 15e; clone k9-1; probe: vaccine; ds.
XX XX
OS Pan troglodytes.
XX XX
XX GB2212511-A.
XX XX
PD 26-JUL-1989.
XX XX
XX 18-NOV-1988; 88GB-0027024.
XX XX

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18-NOV-1987; 87US-0122714.
30-DEC-1987; 87US-0139886.
26-FEB-1986; 88US-0161072.
26-OCT-1988; 88US-0263584.
XX
PA (CHIR) CHIRON CORPORATION.
XX
P2 Houghton M, Choo QL, Kuo G;
XX
XX WPI; 1989-215054/30.
DR P-PSDB; AAP90288.
XX
PT Hepatitis C virus gene - used for prodn. of polynucleotide probes,
PT polypeptide(s) and antibodies for diagnosis, prevention and treatment
PT of infection.
XX
XX Disclosure: fig 47; 235pp; English.
XX
XX The sequence shows a composite hepatitis C virus (HCV) cDNA, derived by
CC aligning clones k9-1 through 15e in 5'-3' direction. The cDNA
CC encodes antigens which react with antibodies in patients with non-A
CC non-B hepatitis (NANBH). The cDNA can be used to design probes, or to
CC synthesize polypeptides, which are used to diagnose HCV-induced NANBH,
CC to raise antibodies for immunoassay or treatment, or to produce
CC vaccines. See also AAP90288, and AAN90303-35.
CC (N.B. This record was resubmitted to correct errors in the sequence.)
CC (Updated on 25-MAR-2003 to correct PR field.)
XX
SQ Sequence 7310 BP; 1495 A; 2216 C; 2058 G; 1539 T; 0 other;

Alignment Scores:
Pred. No.: 1,32e-77 Length: 7310
Score: 943.00 Matches: 179
Percent Similarity: 100.00% Conservatives: 3
Best Local Similarity: 98.35% Mismatches: 0
Query Match: 98.95% Indels: 0
DB: 10 Gaps: 0

US-09-965-594-1 (1-182) x AAN90336 (1-7310)

QY 1 MetAlaProIleThrAlaTyrAlaGlnThrArgGlyLeuLeuGlyCysIleIleThr 20
DB 1728 CTGGCCGCCATCAGCGGTACGCCAGCAGCAGCAAGGGGCTCTAGGGTGCATATCACC 1787

QY 21 SerLeuThrGlyArgAspLysAsnGlnValGluValGlnIleValSerThrAla 40
DB 1788 AGCCTAACTGGCGGACAAAACCAAGTGGAGGGTCAGTTCAGATTTGTGCAACTGCT 1847

QY 41 AlaGlnThrPheLeuAlaThrCysIleAsnGlyValCysTrpThrValTyrHisGlyAla 60
DB 1848 GCCCAACCTTCTGGCAACGTGCATCAATGGGG:GTGGTGGACTGTCTACCCAGCGGGC 1907

QY 61 GlyThrArgThrIleAlaSerProLysGlyProValIleGlnMetTyrThrAsnValAsp 80
DB 1908 GGAACGAGGACCATCGCTCACCAAGGTCCGTGTCATCCAGATGTATACCAATGTAGAC 1967

QY 81 LysAspLeuValGlyTrpProAlaProGlnGlySerArgSerLeuThrProCysThrCys 100
DB 1968 CAAGACCTTGTGGGTGGCGCGCTCCGCAAGGTAGCGGTGTCATTCACACCTGCATTCG 2027

QY 101 GlySerSerAspLeuTyrLeuValThrArgHisAlaAspValIleProValArgAtg 120
DB 2028 GGCCTCTCGSACCTTTACCTGGTCAGGCGACGGGATGTATTCCTCCGTGGCGCGCG 2087

QY 121 GlyAspSerArgGlySerLeuLeuSerProArgProIleSerTyrLeuLysGlySerSer 140
DB 2088 GGTGATAGCAGGGCAGCCTGCTGTCCGCCGCCCATTTCTTACTTTGAAGGCTCCTCG 2147

QY 141 GlyGlyProLeuLeuCysProAlaGlyHisAlaValGlyIlePheArgAlaValCys 160
DB 2148 GGGGGTCCGCTGTGTGTGGCGGGGACCGCGGTGGCATATTTAGGGCGCGGTGTGC 2207

QY 161 ThrArgGlyValAlaLysAlaValAspPheIleProValSerLeuGlnThrMet 180
DB 2208 ACCGTGGAGTGGTAGGGCGGTGGACTTTATCTGTCTGTGAGAACTTAGACAACCATG 2267

QY 181 ArgSer 182
DB 2268 AGCTCC 2273

RESULT 8
AAN90336
ID AAN90336 standard: DNA: 7310 BP.
XX AC AAN90336;
XX AC
XX 25-MAR-2003 (updated)
DT 19-JUL-2001 (updated)
DT 01-NOV-1989 (first entry)
XX XX
DE Composite hepatitis C virus (HCV) cDNA.
KW Hepatitis C virus: cDNA: clone 15e; clone k9-1; probe: vaccine; ds.
XX XX
OS Pan troglodytes.
XX XX
XX GB2212511-A.
XX XX
PD 26-JUL-1989.
XX XX
XX 18-NOV-1988; 88GB-0027024.
XX XX

```

|||||
2208 ACCCGTCAGTGGCTAAGCGGCTGGACTTTATCCCTGTCCAGACCTAGACAGAACCATG 2267
|||||
181 ArgSer 182
2268 AGGTCC 2273

RESULT 9
AAC98221
ID AAC98221 standard; cDNA to mRNA; 7310 BP.
AC AAC98221;
XX
XX
XX 25-MAR-2003 (updated)
XX 15-AUG-1996 (first entry)
XX
DE Hepatitis C virus clone genome.
XX
XX Hepatitis C virus; HCV; antigen: detection; diagnosis; vaccine;
KW antibodies; immunoprophylaxis; sera; serum; ds.
XX
OS Hepatitis C virus.
XX
XX US5443965-A.
XX
XX 22-AUG-1995.
XX
XX 05-APR-1991; 9AUS-0681703.
XX
XX 06-APR-1990; 9OUS-0505621.
XX 09-OCT-1990; 950S-0594854.
XX
XX (GENE-) GENELABS INC.
XX
XX Kim JP, Moekli R, Reyes GR;
XX
XX WPI; 1995-302120/39.
XX
XX New nucleic acids encoding hepatitis C virus antigens - used to
XX develop prods. for detection of hcv-infected sera and prodn. of
XX vaccines and anti-HCV antibodies.
XX
XX Example 4; Figure 11; 71pp; English.
XX
XX Hepatitis C virus (HCV) antigens can be used for detecting HCV
XX infected sera and individuals infected with HCV. They can also be
XX used in an anti-HCV vaccine or for the production of anti-HCV
XX antibodies which can be used for passive immunoprophylaxis. The
XX antigens consistently identify more HCV positive serum samples with
XX a high degree of specificity. See AAC96202-14 and AAR8193-51.
XX (updated on 25-MAR-2003 to correct PF field.)
XX (updated on 25-MAR-2003 to correct PR field.)
XX
XX Sequence 7310 BP; 1454 A; 2217 C; 2060 G; 1589 T; 0 other;

Alignment Scores:
Pred. No.: 132e-77 Length: 7310
Score: 943.60 Matches: 179
Percent Similarity: 100.00% Conservatives: 3
Best Local Similarity: 98.35% Mismatches: 0
Query Match: 98.93% Indels: 0
DB: 16 Gaps: 0

US-09-965-594-1 (1-182) x AAC98221 (1-7310)

QY 1 MetAlaProIleThrAlaIleGlnGlnGlnThrArgGlyLeuLeuGlyCysIleIleThr 20
|||||
DB 1728 CTGGCGCCCATCACCGGTACCGCCAGCAGACAAAGGGCGCTCTAGGTGCATATACCC 1787
|||||
QY 21 SerLeuThrGlyArgAspLysAsnGlyValGluGlyGluValGlnIleValSerThrAla 40
|||||
DB 1788 AGCCTAACTGGCGGGGACAAAAACCAAGTGGAGGGTGAGTCCAGATGTGTCAACTGCT 1847
|||||

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QY 41 AlaGlnThrPheLeuAlaThrCysIleAsnGlyValCysIleThrValIleHisGlyAla 60
|||||
DB 1848 GCCCAACCTTCCTGGCAAGCTGCATCAATGGGGTGTGTGGACTGTCTACCAAGGGGCC 1907
|||||
QY 61 GlyThrArgThrIleAlaSerProLysGlyProValIleGlnMetIleThrAsnValAsp 80
|||||
DB 1908 GGAACGAGGAGACCATGCGTCAOCCAAAGGCTCTGTATCCAGATGTATACCAATGTAGAC 1967
|||||
QY 81 LysAspIleValGlyTrpProAlaProGlnGlySerArgSerLeuThrProCysThrCys 100
|||||
DB 1968 CAAGACCTTGTGGCTGGCGCGCTCGCAAGTACCGCTCATTTGACAGCTGCATTCG 2027
|||||
QY 101 GlySerSerAspLeuTrpLeuValThrArgHisAlaAspValIleProValArgArgArg 120
|||||
DB 2028 GGCTCTCGGACCTTACCTGGTTCAGAGCAGCAGCGCATGTCATTCCTCGTGGCGGCGG 2087
|||||
QY 121 GlyAspSerArgGlySerLeuLeuSerProArgProIleSerTrpLeuLysGlySerSer 140
|||||
DB 2088 GGTGATAGCAGGGGCGACCTGCTGTGCGCGCGCGCCCAITTCCTACTTGAAGGCTCTCG 2147
|||||
QY 141 GlyGlyProLeuLeuCysProAlaGlyHisAlaValGlyIlePheArgAlaAlaValCys 160
|||||
DB 2148 GGGGTCCTGCTGTGTGCGCGCGCGGACGCGCTGGGCATATTTAGGGCGCGGTGTGC 2207
|||||
QY 161 ThrArgGlyValAlaLysAlaValAspPheIleProValIleGluSerLeuIleThrMet 180
|||||
DB 2208 ACCCGTGGAGTGGCTAAGCGGTGGACTTTATCCCTGTGGAGAACCTAGAGACAACCATG 2267
|||||
QY 181 ArgSer 182
|||||
DB 2268 AGGTCC 2273

RESULT 10
AAAT5296
ID AAAT5296 standard; cDNA; 8316 BP.
XX
XX AAAT5296;
XX
XX 15-JAN-2001 (first entry)
XX
XX cDNA sequence compiled Hepatitis C virus cDNA clones.
XX
XX Hepatitis C virus; HCV; antisense polynucleotide; polypeptide;
XX viral infectivity; viral replication; ds.
XX
XX Hepatitis C virus.
XX
XX Key Location/Qualifiers
XX CDS 1..8316
XX /*tag= a
XX /*note= "partial sequence; no termination codon given"
XX
XX EP1034785-A2.
XX
XX 13-SEP-2000.
XX
XX 16-MAR-1990; 2000EP-0109602.
XX
XX 17-MAR-1989; 89US-0325338.
XX 20-APR-1989; 89US-0341334.
XX 18-MAY-1989; 89US-0355002.
XX 16-MAR-1990; 90EP-0302866.
XX
XX (CHIR ) CHIRON CORP.
XX
XX Houghton M, Choo Q, Kuo G;
XX
XX WPI; 2000-566897/53.
XX P-PSDB; AAB18540.
XX
XX Novel composition comprising a hepatitis C virus antisense
XX polynucleotide which is complementary to or corresponds to a sense
XX strand of the virus genome, and selectively hybridises to it.
XX
XX

```

```
XX
PS
XX
XX The specification describes a pharmaceutical composition which
CC comprises a hepatitis C virus (HCV) antisense polynucleotide. The
CC HCV is characterized by a positive stranded RNA genome which has
CC 40% homology at the polypeptide level to a HCV polypeptide. The
CC antisense polynucleotide binds to cellular polynucleotides which
CC enhance and/or are required for viral infectivity, replicative
CC ability or chronicity. The antisense polynucleotides may also be
CC designed to bind with high specificity, to be of increased stability,
CC to be stable and to have low toxicity. The composition also comprises
CC an agent which causes viral RNA to be inactive. The composition
CC is used for preventing HCV replication in a system. The present
CC sequence represents a novel HCV cDNA sequence, which is used in the
CC course of the invention.
XX
SQ Sequence 8316 BP; 1671 A; 2529 C; 2345 G; 1771 T; 0 other:
Alignment Scores:
Pred. No.: 1,558-77 Length: 8316
Score: 943.00 Matches: 179
Percent Similarity: 100.00% Conservative: 3
Best Local Similarity: 98.95% Mismatches: 0
Query Match: 98.95% Indels: 0
DB: 21 Gaps: 3
US-09-965-594-1 (1-182) x AA75296 (1-816)
CY : MetAlaPcTtLeThrAlaTyraAlaGlnGlnThrArgGlyGlyGlnGlyCysIleIleCpT 20
DB 2734 CTGGGGGCGGATCACCGGCTACCGCCGACAGACAGCGCCCTAGGCTGATATACCC 2793
CY 21 SerLeuThrGlyArgAspLysAsnGlnValGlnGlyGlnValGlnIleValSerThrAla 40
DB 2794 AGCCTAACTGGCGGGGACAAAACCAAGTGAAGGTAGGTCAGATTGCTCAACTGCT 2853
CY 41 AlaGlnThrPheLeuAlaThrCysIleAsnGlyValCysThrPheValThrHisLeuAla 60
DB 2854 GCCCAAACTTCCTGGCAACGTGCACTCAATGAGGTGCTCTGGACGTGCTACGACGGGACC 2913
CY 61 GlyThrArgThrIleAlaSerProLysGlyProValIleGlnMetTyrThrAspValAsp 80
DB 2914 CGAAGGAGGACCATCGCGCACCAAGGCTCTGTGAACAGATGTATACCAAGIAGAC 2973
CY 81 LysAspLeuValGlyTyrProAlaProGlnGlySerArgSerLeuThrProCysThrCys 100
DB 2974 CAAGACCTTGCGGCTGCGCTCGCAAGGTAGCCGCTCATGAGACCGCTGCACCTTC 3033
CY 101 GlySerSerAspLeuTyrLeuValThrArgHisAlaAspValIleProValArgArgArg 120
DB 3034 GGTCTCTGGACCTTATCTGTCTCAGGAGGACGCGGATGTCTTCCTGGTGGCGGGGAG 3093
CY 121 GlyAspSerArgGlySerLeuLeuSerProArgProIleSerTyrLeuLysGlySerSer 140
DB 3094 GTGTATAGAGGGGAGCGCTGTGTGCGCGCGCGGCGCATTTCCACTTGAAGGCTCCICG 3153
CY 141 GlyGlyProLeuLeuCysProAlaGlyHisAlaValGlyIlePheArgAlaAlaValCys 160
DB 3154 GGGGGTCCGCTGTGTGCGCGCGCGGACCGCGCTGGGCATATTAGGCGCGCGGTGCG 3213
CY 161 ThrArgGlyValAlaLysAlaValAspPheIleProValGluSerLeuLeuIleThrMet 180
DB 3214 ACCCGTGGAGTGCTAAGCGGTGGACTTTATCCCTGGGAGAGACTAGACAAACCATG 3273
CY 181 ArgSer 182
DB 3274 AGGTCC 3279
RESULT 11
AAZ07656
ID AAZ07656 standard; DNA: 9133 BP.
XX
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AC
XX
XX AA207656;
DI 20-MAR-2003 (updated)
XX 08-NOV-1999 (first entry)
XX Nucleotide sequence of HCV-1 ORF.
XX Hepatitis C virus; HCV; J1; J7; HCV-1; non-A, non-B HCV; NANBH;
XX HCV infection; vaccine; ds.
XX Hepatitis C virus.
XX Key Location/Qualifiers
XX CDS 268..9132
XX /*tag=
XX /transl_except= (pos:1588..1589; aa:Leu)
XX /note= "this codon has an apparent 1 nucleotide deletion,
XX which alters the reading frame"
XX /transl_except= (pos:1647..1650; aa:Pro)
XX /note= "this codon has an apparent 1 nucleotide
XX insertion, which alters the reading frame; this
XX insertion is not indicated in the sequence
XX present in the formal sequence listing of the
XX specification"
XX
XX EP939128-A2.
XX
XX C1-SEP-1999.
XX
XX 17-SEP-1990; 99EP-0101746.
XX
XX 15-SEP-1989; 89CS-0408045.
XX 21-DEC-1989; 89CS-0436142.
XX 17-SEP-1990; 90EP-C3-0149.
XX
XX (CHIR ) CHIRON CORP.
XX (CYAAV) OYA A.
XX
XX Cha T, Han J, Houghton M, Irvine BD, Kolberg JA;
XX Miyamura T, Saito I, Weiner AJ;
XX
XX WPI: 1599-460643/41.
XX P-PSDB: AAY:4375.
XX
XX New Hepatitis C Virus isolates, useful for diagnosis of hepatitis
XX infections and development of vaccines
XX
XX Disclosure: Fig 12; 132pp; English.
XX
XX The invention provides two new isolates of hepatitis C virus (HCV). J1
XX and J7. These two isolates comprise nucleotide and amino acid sequences
XX that are distinct from the HCV isolate HCV-1. The nucleotide sequences
XX may be used to detect non-A, non-B HCV (NANBH) polynucleotides by
XX hybridisation for diagnosis of NANBH infections. They may also be used to
XX screen blood donors, donated blood and blood products for this infection.
XX The isolates may also be used to isolate other naturally occurring
XX variants of the virus. The polypeptides may be used as a vaccine for
XX administration to patients to protect against infection with NANBH. The
XX present sequence represents the nucleotide sequence of HCV-1 ORF.
XX (Updated on 20-MAR-2003 to correct PF field.)
XX (Updated on 20-MAR-2003 to correct PR field.)
XX
XX Sequence 9133 BP; 1834 A; 2772 C; 2600 G; 1927 T; 0 other:
Alignment Scores:
Pred. No.: 1,748-77 Length: 9133
Score: 943.00 Matches: 179
Percent Similarity: 100.00% Conservative: 3
Best Local Similarity: 98.95% Mismatches: 0
Query Match: 98.95% Indels: 0
DB: 20 Gaps: 0
US-09-965-594-1 (1-182) x AAZ07656 (1-9133)
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QY      1 MetAlaProIleThrAlaTyrAlaGlnGlnThrArgGlyLeuLeuGlyCysIleIleThr 20
DB      3343 CTGGCGCCATCAGCGGCTAGCCACACACAGAGGCTCTCTAGAGTCATAACACACC 3402
QY      21 SerLeuThrGlyArgAspLysAsnGlnValGluGlyGluValGlnIleValSerThrAla 40
DB      3403 AGCCTAACTGGCCGGGACAAAACCAAGTGCAGGGTCAGAGTGTCTCAACTGCT 3462
QY      41 AlaGlnThrPheLeuAlaThrCysIleAsnGlyValCysIleThrValTyrHisGlyVala 60
DB      3463 GCCCAACCTTCCTGGCAAGCTGCATCAATGGGCTGTCGAGCTGTACCCAGGGGCC 3522
QY      61 GlyThrArgThrIleAlaSerProLysGlyProValIleGlnMetTyrThrAsnValAsp 80
DB      3523 GGAAGAGGAGGACCATCGKGTCCACCAAGGTCCTCTCATCCAGATGATACCAATGTAGAC 3582
QY      81 LysAspLeuValGlyTyrProAlaProGlnGlySerArgSerLeuThrProCysThrCys 100
DB      3583 CAAGACCTTGTGGCTGGCCGCTCCGCAAGGAGCGGCTCATTCACACCTGCCTTGC 3642
QY      101 GlySerSerAspLeuTyrLeuValThrArgHisAlaAspValIleProValArgArg 120
DB      3643 GGCTCCCTCGACCTTACCTGGTCACAGAGCAGCCGCAATGCTATCCCTGCGCGCGCG 3702
QY      121 GlyAspSerArgGlySerLeuLeuSerProArgProIleSerTyrLeuLysGlySerSer 140
DB      3703 GGTGATAGCAGGGCAGCGCTGCTGCGCGCGCGCCATTTCTACTTGAAAGGCTCTCG 3762
QY      141 GlyGlyProLeuLeuCysProAlaGlyHisAlaValGlyIlePheArgAlaAlaValCys 160
DB      3763 GGGGTCGCGCTGTGTGCGCGCGCGGCGCGCGCGCGCGCATATTTAGGGCGCGGTGTC 3822
QY      161 ThrArgGlyValAlaLysAlaValAspPheIleProValGluSerLeuGluThrThrMet 180
DB      3823 ACCCGTGGAGTGGTAAGCGGCTTGATTCCTTATCCCTGTGGAGACCTAGACAAACCA 3882
QY      181 ArgSer 182
DB      3883 AGGTCC 3888

RESULT 12
ID      AAQ05956 standard; DNA; 9185 bp.
AC      AAQ05956;
DT      25-MAR-2003 (updated)
DT      23-JAN-1991 (first entry)
XX      Sense strand of the compiled Hepatitis C virus cDNA sequence.
DE      Hepatitis C virus (HCV); antiviral agent; ss.
KW      Hepatitis C virus.
OS      Hepatitis C virus.
XX      Key Location/Qualifiers
FT      CDS 320..9185
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FT      misc_RNA 1..1667
FT      /*tag= b
FT      /*note="epitope within this region is claimed"
FT      misc_RNA 8978..9185
FT      /*tag= c
FT      /*note="encodes an epitope that is claimed"
XX      EP368232-A.
XX      19-SEP-1990.
XX      16-MAR-1990; 9CEP-0302665.
XX      18-MAY-1989; 89US-0355002.

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PR      17-MAR-1989; 89US-0325338.
PR      20-APR-1989; 89US-0341334.
XX      (CHIR ) CHIRON CORP.
XX      Houghton M, Choo QL, Kuo G.
XX      WPI: 1990-284416/38.
XX      P-PSDB; AAR08124.
XX      Hepatitis C virus DNA - used for producing probes,
XX      polypeptide(s), antibodies and anti-sense polynucleotide(s) for
XX      diagnosis and therapy.
XX      Disclosure: Fig 17; 83pp; English.
XX      HCV cDNA libraries were constructed using pooled serum from a
XX      chimpanzee with chronic HCV infection. A lambda gt10 library was
XX      screened with probes derived from previously isolated clones. The
XX      ORF is derived from the overlapping clones b14a, aq30a, CA205a,
XX      CA290a, CA216a, p14a, CA167b, CA156a, CA59a, K9-1, 253, 131,
XX      121, 141, 11b, 7f, 8h, 33c, 40b, 37b, 35, 36, 81, 32, 33b, 25c,
XX      14c, 8f, 33f, 33d, 39c, 35f, 19g, 26g, 15e, b5a and 16jh. These
XX      clones extend the sequence of the HCV genome reported in EP-38216.
XX      The upstream region from nucleotides -319 to +1348 (=11667 in this
XX      file) is covered by clones b14a, 18g, aq30a, CA205a, CA250a,
XX      CA216a, p14a, CA167b, CA156a, CA84a and CA59a; nucleotides
XX      8559-8866 (=8978-9185 in this file) are covered by clones b5a and
XX      16jh.
XX      See also AAQ05955.
XX      CC (Updated on 25-MAR-2003 to correct PA field.)
XX      SQ Sequence 9185 BP; 1849 A; 2790 C; 2608 G; 1938 T; 0 other;

Alignment Scores:
Pred. No.: 1,750-77 Length: 9185
Score: 943.00 Matches: 179
Percent Similarity: 100.00% Conservative: 3
Best Local Similarity: 98.35% Mismatches: 0
Query Match: 98.95% Indels: 0
DB: 11 Gaps: 0

US-09-965-594-1 (1-182) x AAQ05956 (1-9185)
QY      1 MetAlaProIleThrAlaTyrAlaGlnGlnThrArgGlyLeuLeuGlyCysIleIleThr 20
DB      3395 CTGGCGCCATCAGCGGCTAGCCACACACAGAGGCTCTCTAGAGTCATAACACACC 3454
QY      21 SerLeuThrGlyArgAspLysAsnGlnValGluGlyGluValGlnIleValSerThrAla 40
DB      3455 AGCCTAACTGGCGGACAAAACCAAGTGCAGGGTCAGGTCCAGATTGTGTCAACTGCT 3514
QY      41 AlaGlnThrPheLeuAlaThrCysIleAsnGlyValCysIleThrValTyrHisGlyVala 60
DB      3515 GCCCAACCTTCCTGGCAAGCTGCATCAATGGGCTGTCTGAGACTGTCTACACGGGGCC 3574
QY      61 GlyThrArgThrIleAlaSerProLysGlyProValIleGlnMetTyrThrAsnValAsp 80
DB      3575 GGAACGAGGACCATCGCTCACCACAGGGTCCTGTATCCAGATGTATACCAATGTAGAC 3634
QY      81 LysAspLeuValGlyTyrProAlaProGlnGlySerArgSerLeuThrProCysThrCys 100
DB      3635 CAAGACCTTGTGGCTGGCCGCTCCGCAAGGTAGCGGCTCATTCACACCTGCACCTTGC 3694
QY      101 GlySerSerAspLeuTyrLeuValThrArgHisAlaAspValIleProValArgArg 120
DB      3695 GGCTCCCTCGACCTTACCTGGTCAGAGGCGAGCGGATGTCAITCCCTGGCGCGCGG 3754
QY      121 GlyAspSerArgGlySerLeuLeuSerProArgProIleSerTyrLeuLysGlySerSer 140
DB      3755 GGTGATAGCAGGGCAGCGCTGTGTCGCGCGCGCGCATTTCTACTTGAAGGCTCTCTCG 3814
QY      141 GlyGlyProLeuLeuCysProAlaGlyHisAlaValGlyIlePheArgAlaAlaValCys 160

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Db 3815 GGGGTCCCGTGTGTGCGCGGGGACGCGGTGGGCATATTAGGCGCGGGTGTGC 3874
Qy 161 ThrArgGlyValAlaLysAlaValAspPheIleProValGluSerLeuGluThrMet 180
Db 3875 ACCGTGAGTGGCTAAGCGGTGGACTTATCCGTGGAGAACCTAGACACCACTG 3934
Qy 181 ArgSer 182
Db 3935 AGGTCC 3940

RESULT 13
ID AAQ10566 standard; DNA; 9185 BP.
XX
AC AAQ10566:
DT 25-MAR-2003 (updated)
DT 29-APR-1991 (first entry)
XX
DE Hepatitis C virus strain 1 DNA.
XX
KW Hepatitis C virus; HCV-1; non-A, non-B hepatitis; HCV antigen;
KW viral infections; ss.
XX
OS Hepatitis C virus.
XX
PN EP414475-A.
XX
PD 27-FEB-1991.
XX
PF 21-AUG-1990; 90FP-0309120.
XX
PR 25-AUG-1989; 89US-0398667.
XX
PA (CHTR ) CHIRON CORP.
XX
PI Weiner AJ, Steimer KS;
XX
DR WPI; 1991-059670/79.
XX
PT Cell lines infected with hepatitis C virus - are used as source
PT of antigens for detection of HCV antibodies, for vaccines, and
PT for screening anti-viral agents
XX
PS Disclosure; fig 1; 24pp; English.
XX
CC This is a hepatitis C virus (HCV) composite cDNA sequence, deduced
CC using overlapping clones, a compo. contg. the antigenic protein
CC encoded by this sequence is useful for detecting anti-HCV anti-
CC bodies (Abs) and for screening an agent which inhibits HCV repli-
CC cation. A cell line infected with this virus can be used as a
CC source of antigens. The antigen is useful for preparing vaccines
CC for treating viral infections. See also AAQ10567.
CC (Updated on 25-MAR-2003 to correct PA field.)
XX
SQ Sequence 9185 BP; 1845 A; 2790 C; 2603 G; 1938 T; 0 other;

Alignment Scores:
Pred. No.: 1.75e-77 Length: 9185
Score: 943.00 Matches: 179
Percent Similarity: 100.00% Conservative: 3
Best Local Similarity: 98.35% Mismatches: 0
Query Match: 98.95% Indels: 0
DB: 12 Gaps: 0

US-09-965-594-1 (-182) x AAQ10566 (1-9185)

Qy 1 MetAlaProIleThrAlaValAlaGlnGlnThrArgGlyLeuLeuGlyCysIleIleThr 20
Db 3395 CTGGCGCCCATCACGCGGTACGCCAGACAAAGGGCGCTCTAGGGTGCATAATCACC 3454
Qy 21 SerLeuThrGlyArgAspLysAsnGlnValGluGlyGluValGlnIleValSerThrAla 40

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Db 3455 AGCCTAACTGCGCGGACAAAACCAAGTGCAGGGTGAGGTCCAGATTGTCTCAACTGCT 3514
Qy 41 AlaGlnThrPheLeuAlaThrCysIleAsnGlyValCysTrpIleValIleHisGlyAla 60
Db 3515 GCCCAACCTTCCTGGCAACGTGCATCAATGGGGTGTGCTGGAGTGTCTACACGGGGCC 3574
Qy 61 GlyThrArgThrIleAlaSerProIysGlyProValIleGlnMetTyrThrAsnValAsp 80
Db 3575 GCAACGAGGACCATCGCTCACCAAGGGTCTGTCTATCCAGATGTATACCAATGTAGAC 3634
Qy 81 LysAspLeuValGlyTrpProAlaProGlnGlySerArgSerLeuThrProCysThrCys 100
Db 3635 CAAGACCTTGTGGCTGGCGCTCCGCAAGGTAGCGGCTCATTGACACACCTGCACCTTGC 3694
Qy 101 GlySerSerAspLeuTyrLeuValThrArgHisAlaAspValIleProValArgArg 120
Db 3695 GGTCTCTGGACCTTTACCTGGTCACGAGGACCGCGATGTCATCCCGTGGCGGCGG 3754
Qy 121 GlyAspSerArgGlySerLeuLeuSerProArgProIleSerTyrLeuLysGlySerSer 140
Db 3755 GGTATTAGCAGGGGACGCTGCTGCCCGCGGCCATTCTCTACTTCAAAGGCTCCTCG 3814
Qy 141 GlyGlyProLeuLeuCysProAlaGlyHisAlaValGlyIlePheArgAlaAlaValCys 160
Db 3815 GGGGTCGGCTGTGTGCGCGCGGACCGCGTGGGCATATTAGGCGCGCGGTGTGC 3874
Qy 151 ThrArgGlyValAlaLysAlaValAspPheIleProValGluSerLeuGluThrMet 180
Db 3875 ACCGTGAGTGGCTAAGCGGTGGACTTATCCGTGGAGAACCTAGACACCACTG 3934
Qy 181 ArgSer 182
Db 3935 AGGTCC 3940

RESULT 14
ID AAA75297 standard; cDNA; 9185 BP.
XX
AC AAA75297:
XX
DT 15-JAN-2001 (first entry)
XX
DE Sense strand of HCV encoding a polyprotein.
XX
KW Hepatitis C virus; HCV; antisense polynucleotide; polyprotein;
KW viral infectivity; viral replication; ds.
XX
OS Hepatitis C virus.
XX
FH Key Location/Qualifiers
FT CDS 320..9184
FT /*tag= a
FT /note= "partial sequence; no termination codon given"
XX
PN EP3034785-A2.
XX
PD 13-SEP-2000.
XX
PE 16-MAR-1990; 2000EP-0109602.
XX
PR 17-MAR-1989; 89US-0325338.
XX
PR 20-APR-1989; 89US-0341334.
XX
PR 18-MAY-1989; 89US-0355002.
XX
PR 16-MAR-1990; 90EP-0302866.
XX
PA (CHTR ) CHIRON CORP.
XX
PI Houghton M, Choo Q, Kuo G;
XX
DR WPI; 2000-566891/53.
XX
DR P-PSDB; AAB18541.

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GenCore version 5.1.6
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OM protein - nucleic search, using frame_plus_p2n model

Run on: September 27, 2003, 12:22:50 : Search time 3348 Seconds
(without alignments)
2223.881 Million cell updates/sec

Title: US-09-965-594-1

Perfect score: 953

Sequence: : MAP::YAAQTRGL:GCTIT.....GVAKAVDFIPVSLDT:MRS 182

Scoring table: BLOSUM62

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Ygapop 10.0 , Ygapext 0.5

Fgapop 6.0 , Fgapext 7.0

Dgapop 6.0 , Dgapext 7.0

Searched: 2668711 seqs, 2045483396 residues

Total number of hits satisfying chosen parameters: 577422

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 99%

Listing first 45 summaries

Command line parameters:

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-DSV_TIMEOUT=120 -WARN_TIMEOUT=30 -THREAFS=1 -XGAPOP=10 -XGAPEXT=0.5 -Fgapop=6
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Database :

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2: gb_hg:

3: gb_in:

4: gb_om:

5: gb_ov:

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9: gb_pr:

10: gb_ro:

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14: gb_vl:

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18: em_in:

19: em_mu:

20: em_om:

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23: em_pat:

24: em_ph:

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match %	Length	DB ID	Description
1	946	99.3	2058	6	AX395309 Sequence
2	946	99.3	2058	6	AX454818 Sequence
3	945	99.2	543	14	AF369218 Hepatitis
4	945	99.2	543	14	AF369235 Hepatitis
5	943	99.0	5360	6	AR118686 Sequence
6	943	99.0	5360	6	I06434 Sequence 48
7	943	99.0	5360	6	I03328 Sequence 8
8	943	99.0	6785	6	AR118692 Sequence
9	943	99.0	6785	6	I06440 Sequence 54
10	943	99.0	6785	6	I03329 Sequence 10
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14	943	99.0	8316	6	AR118703 Sequence
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16	943	99.0	9185	6	AR118722 Sequence
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19	943	99.0	9185	5	I08254 Sequence 1
20	943	99.0	9379	6	AR166930 Sequence
21	943	99.0	9379	6	AR301300 Sequence
22	943	99.0	9401	6	AR176483 Sequence
23	943	99.0	9401	6	BD080334 Hepatitis
24	943	99.0	9401	6	E65593 Hepatitis C
25	943	99.0	9401	6	I71894 Sequence 9
26	943	99.0	9401	6	I81885 Sequence 9
27	943	99.0	9401	14	HPCPLYPRE M62321 Hepatitis C
28	943	99.0	9409	12	AF387805 Synthetic
29	943	99.0	9409	12	AF387808 Synthetic
30	943	99.0	9418	14	AF274632 Hepatitis
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32	943	99.0	9693	12	AF387807 Synthetic
33	942	98.8	543	14	AF369222 Hepatitis
34	942	98.8	543	14	AF369232 Hepatitis
35	942	98.8	543	14	AF369240 Hepatitis
36	942	98.8	543	14	AF369245 Hepatitis
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38	942	98.8	2061	6	AX457113 Sequence
39	940	98.6	543	14	AF369224 Hepatitis
40	940	98.6	543	14	AF369237 Hepatitis
41	940	98.6	9424	14	AF511948 Hepatitis
42	939	98.5	543	14	AF369230 Hepatitis
43	939	98.5	9502	6	E08263 gRNA of Hep
44	939	98.5	9502	6	E08264 cDNA of Hep
45	939	98.5	9502	14	HFCHCJ1 D10749 Hepatitis C

ALIGNMENTS

RESULT 1

AX395309	AX395309	2058 bp	DNA	linear	PAT 18-MAY-2003
LOCUS	Sequence 2 from Patent WO0196875.				
DEFINITION	AX395309				
ACCESSION	AX395309				
VERSION	AX395309.1	GI:21066308			
KEYWORDS					
SOURCE	synthetic construct				
ORGANISM	artificial sequences.				
REFERENCE	1				
AUTHORS	Chier,D.V., Arcangei,P., Iandreske,L., George-Nasole,ent),C., Colt,D. and Medina-Seiby,A. Ncr antigen/body combination assay Patent: WO 0196875-A 2 28-DEC-2001 CHIRON CORPORATION (US) Location/Organism: 1..2058 /organism "Synthetic Construct" /mol_type "genomic DNA" /db_xref "taxon:32630" /notes "representative KSE/4a conformational antigen"				
TITLE					
JOURNAL					
FEATURES					
SOURCE					
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ORIGIN					
Alignment Scores:					
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Qy	21 SerLeuThrCiyArgAspIysAsnGlnValLLeGlyGluValIleIleValSerThraLa	40			
Db	61 AGCTTAACCTGGCGGGACAAAACCAACTGGAGGCTGANGTCCAGATTGATCAACTGCI	120			
Qy	41 AlaGlnThrPheLeuAlaThrCysIleAsnLysValCysTrpThrValTryHisGlyVal	60			
Db	121 GCCCAACCTTCTTGGCAACGTGCATCAATACGGGTGCTGTGSA-TGCTACCCACGGGCC	180			
Qy	61 GlyThrArgThrIcAlaSerProIysGlyProValIleGlnMetTyThrAsnValasp	80			
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Qy	31 LysAspLeuValGlyTrpProAlaProCingLysSerArqSerLeuThrProCysThrCys	100			
Db	241 CAAGACCTTCTGGCTGGCCGGTCCCAAGGTAGCGGATCATTTGACACCTGACCTTGC	300			

Qy	101	GlySerSerAspLeuTyrLeuValThrArgHisAlaAspValIleProValArgArg	120		
Db	301	GGCTCTCGGACCTTTACCTGGTGCACGAGCACCGCGATGTCATTC	360		
Qy	121	GlyAspSerArgGlySerLeuLeuSerProArcProIleSerTyrLeuIysGlySerSer	140		
Db	361	GGTGATAGAGGGGAGCGCTGCTGCGCCCGCGCCATATTTCCACTTGAAGGCTCCTCG	420		
Qy	141	GlyGlyProLeuLeuLeuCysProAagIyHsAlaValAlaGlyIlePheArgAlaAlaValCys	160		
Db	421	GGSGGTCCGCTGTTGTCGCCCGGGCGACGCGGCGGCATATTTAGGGCGCGGTGTGC	480		
Qy	161	ThrArgGlyValAlaIysAlaValAlaAspPheIleProValIleSerLeuGluThrMet	180		
Db	481	ACCGGTGGAGTGCTAAGCGGTGGACTTTATCCCTGTGGAGAACTAGACAGCAACCATG	540		
Qy	181	ArgSer	182		
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RESULT 2					
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LOCUS					
DEFINITION		Sequence 1 from Patent WO0196870.			
ACCESSION		AX454818			
VERSION		AX454818.1			
KEYWORDS		GI:21714047			
SOURCE		Synthetic construct			
ORGANISM		Synthetic construct			
REFERENCE		artificial sequences.			
AUTHORS		Chien,D.Y., Arcangel,P., Tandeske,L., George-Nascimento,C.,			
TITLE		Coit,D. and Medina-Solby,A.			
JOURNAL		Immunassays for anti-hcv antibodies			
		Patent: WO 0196870-A: 1 20-DEC-2001:			
		CHIRON CORPORATION (US)			
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		SDLYVTRADVPIVPRRRDGRSLSPRISTYLGKSGGPLLCPAGHVGIFRAVAC			
		TRGVAKAVDFIVENLETMRSPFTDNSPPVQSFQVHLHAFPTGSKSTKPLA			
		YAAQGYKVLVLPNSVAATFGQAYMSKAHGIQDPIRITVRIITIGSPIITYSGFLA			
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		PHIEEVALSTGEIPEYKVAIDALMTGYTGDFSDVIDCNTCVTVDFSLDPTFTLT			
		YKGLDVPITPGDVVVVATDTPGTVREVPGERSPGMSFVCECYDAGCANVELTPA			
		TLTPQDAVSTORRGIRGKPKGYREVPGERSPGMSFVCECYDAGCANVELTPA			
		ETVRURATMTNFTGLPCVDHPIFWESGVTIGLIDAHFSLQTKSGENLPLVATQA			
		TVCAKAAFPSPNDQMKKCLRLKPLLHGPTPLIYELGAVQNEITLTHPTVTKYIMTCM			
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BASE COUNT	419 a	633 c	581 g	425 t	
ORIGIN					
Alignment Scores:					
Pred. No.:	6,56e-68	Length:	2058		
Score:	946.00	Matches:	180		
Percent Similarity:	100.00%	Conservative:	2		
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Query Match:	99.27%	Indels:	0		
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QY	21	SerLeuThrGlyArgAspIysAsnGlnValGlnGlyGluValGlnIleValSerThrAla	40
DB	61	AGGCTACCTGCGCGGACAAAACCAAGTGGAGGTGAGGTCACATGTCTCAACTGCT	120
QY	41	AlaGlnThrPheLeuAlaThrCysIleAsnGlyValCysIleThrValTyrHisGlyVala	60
DB	121	GCCCAACCTTCTCGCAGCTGCATCAATGAGGTGTGCTGACTGCTTACACAGCGGCT	180
QY	61	GlyThrArgThrIleAlaSerProGlyGlyProValIleGlnMetTyrThrAsnValasp	80
DB	181	GGAACGAGGACCATCGCTGCCCAAGGCTGTGTGTCATCCAGATGTATACCAATGTAGAC	240
QY	81	LysAspLeuValGlyThrProAlaProGlnGlySerArgSerLeuThrProCysThrCys	100
DB	241	CAAGACTTGTGGCTGGCGCGCTCCGCAAGGTAGCGATCATTCACACCTGCACATGC	300
QY	101	GlySerSerAspLeuTyrLeuValThrArgHisAlaAspValIleProValArgArg	120
DB	301	GGCTCTCGGACCTTTCCTGTGTCAGGAGCGACCGGATGTCATTCCTGCGCGCGGG	360
QY	121	GlyAspSerArgGlySerLeuLeuSerProArgProIleSerTyrLeuLysGlySerSer	140
DB	361	GGTCATAGCAGGCGCATCGCTGTGCTGCGCGCGCGCAATTCCTACTGAAAGGCTCT	420
QY	141	GlyGlyProLeuLeuCysProAlaGlyHisAlaValGlyIlePheArgAlaAlaValCys	160
DB	421	GGGGGTCCTGT	480
QY	161	ThrArgGlyValAlaLysAlaValAspPheIleProValGlySerLeuThrThrMet	180
DB	481	ACCGTGGAGTGGCTAAGCGGTGACTTATCTCTGTGGAGAACCTAGACAAACCAATG	540
QY	181	ArgSer 182	
DB	541	AGGTCC 546	
RESULT 3			
LOCUS	AF369218	543 bp RNA linear VRL 03-JUL-2002	
DEFINITION	Hepatitis C virus Pt.4 NS3 protease gene, partial cds.		
ACCESSION	AF369218		
VERSION	AF369218.1	GI:14150550	
KEYWORDS	Hepatitis C virus		
SOURCE	Hepatitis C virus		
ORGANISM	Hepatitis C virus		
REFERENCE			
AUTHORS	Holland-Staley,C.A., Kovari,L.C., Golenberg,E.M., Pekarsky,K.J. and Mayers,D.L.		
TITLE	Genetic diversity and response to IFN of the NS3 protease gene from clinical strains of the hepatitis C virus		
JOURNAL	Arch. Virol. 147 (7), 1385-1406 (2002)		
MEDLINE	22105140		
PUBMED	12111414		
REFERENCE			
AUTHORS	Holland-Staley,C.A., Kovari,L.C., Golenberg,E. and Mayers,D.L.		
TITLE	Genetic diversity and response to IFN of the NS3 protease gene from clinical strains of the hepatitis C virus		
JOURNAL	Arch. Virol. 147 (7), 1385-1406 (2002)		
MEDLINE	22105140		
PUBMED	12111414		
REFERENCE			
AUTHORS	Holland-Staley,C.A., Kovari,L.C., Golenberg,E. and Mayers,D.L.		
TITLE	Genetic diversity and response to IFN of the NS3 protease gene from clinical strains of the hepatitis C virus		
JOURNAL	Arch. Virol. 147 (7), 1385-1406 (2002)		
MEDLINE	22105140		
PUBMED	12111414		
REFERENCE			
AUTHORS	Holland-Staley,C.A., Kovari,L.C., Golenberg,E. and Mayers,D.L.		
TITLE	Genetic diversity and response to IFN of the NS3 protease gene from clinical strains of the hepatitis C virus		
JOURNAL	Arch. Virol. 147 (7), 1385-1406 (2002)		
MEDLINE	22105140		
PUBMED	12111414		
REFERENCE			
AUTHORS	Holland-Staley,C.A., Kovari,L.C., Golenberg,E. and Mayers,D.L.		
TITLE	Genetic diversity and response to IFN of the NS3 protease gene from clinical strains of the hepatitis C virus		
JOURNAL	Arch. Virol. 147 (7), 1385-1406 (2002)		
MEDLINE	22105140		
PUBMED	12111414		
REFERENCE			
AUTHORS	Holland-Staley,C.A., Kovari,L.C., Golenberg,E. and Mayers,D.L.		
TITLE	Genetic diversity and response to IFN of the NS3 protease gene from clinical strains of the hepatitis C virus		
JOURNAL	Arch. Virol. 147 (7), 1385-1406 (2002)		
MEDLINE	22105140		
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REFERENCE			
AUTHORS	Holland-Staley,C.A., Kovari,L.C., Golenberg,E. and Mayers,D.L.		
TITLE	Genetic diversity and response to IFN of the NS3 protease gene from clinical strains of the hepatitis C virus		
JOURNAL	Arch. Virol. 147 (7), 1385-1406 (2002)		
MEDLINE	22105140		
PUBMED	12111414		
REFERENCE			
AUTHORS	Holland-Staley,C.A., Kovari,L.C., Golenberg,E. and Mayers,D.L.		
TITLE	Genetic diversity and response to IFN of the NS3 protease gene from clinical strains of the hepatitis C virus		
JOURNAL	Arch. Virol. 147 (7), 1385-1406 (2002)		
MEDLINE	22105140		
PUBMED	12111414		
REFERENCE			
AUTHORS	Holland-Staley,C.A., Kovari,L.C., Golenberg,E. and Mayers,D.L.		
TITLE	Genetic diversity and response to IFN of the NS3 protease gene from clinical strains of the hepatitis C virus		
JOURNAL	Arch. Virol. 147 (7), 1385-1406 (2002)		
MEDLINE	22105140		
PUBMED	12111414		
REFERENCE			</

QY 161 ThrArgGlyValAlaIysAlaValAspPheIleProValGluSerLeuGluThrThrMet 160
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 Db 1410 ACCCGTGGAGTGGCTAGAGGGGTGGACTTATCCCTGTGGAGAACCTAGAGACACCATG 1469
 QY 161 ArgSer 182
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RESULT 6
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 LOCUS 106434
 DEFINITION Sequence 48 from Patent EP 0318216.
 ACCESSION 106434
 VERSION 106434.1 GI:590311
 KEYWORDS
 SOURCE Unknown.
 ORGANISM Unknown.
 REFERENCE 1 (bases 1 to 5360)
 AUTHORS Houghton,M., Choo,Q.-L. and Kuo,G.
 TITLE Nanbv diagnostics and vaccines
 JOURNAL Patent: EP 0318216-A1 48 31-MAY-1989;
 FEATURES
 location/Qualifiers
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 /organism="unknown"
 BASE COUNT 1061 a 1623 c 1533 g 1143 t
 ORIGIN

Alignment Scores:
 Pred. No.: 3,226-67 Length: 5360
 Score: 943.00 Matches: 179
 Percent Similarity: 100.00% Conservative: 3
 Best Local Similarity: 98.35% Mismatches: 0
 Query Match: 98.95% Indels: 0
 DB: 6 Gaps: 0

US-09-965-594-1 (1-182) x 106434 (1-5360)

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 QY 21 SerLeuThrGlyArgAspLysAsnGlnValGlyGluValGlnIleValSerThrAla 40
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 Db 990 AGCTTAACISGCCGAGACAAACACAGTGAAGGTGAGGTCCAGATTCGTACTGCT 149
 QY 41 AlaGlnThrPheLeuAlaThrCysIleAsnGlyValCysTrpThrValCysHisGlyAla 60
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 Db 1050 GGCACAAACCTTCCTGCAACAGTGCATCAATGGGTGTGTGACTGTCTACCAAGGGGCC 1169
 QY 61 GlyThrArgThrIleAlaSerProLysGlyProValIleGlnMetTyrThrAsnValAsp 80
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 Db 1110 GGAACAGAGACCATCCGTCACCAAGGTCCTGTCTCATCCAGATGTATACCAATGTAGAC 1169
 QY 81 LysAspLeuValGlyTrpProAlaProGlnGlySerArgSerLeuThrProCysThrCys 100
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 Db 1170 CAAGACCTTGTGGGTGGGCCCTCCGCAAGGTAGCCGCTCATTCACACCTGCACTTGC 1229
 QY 101 GlySerSerAspLeuTyrLeuValThrArgHisAlaAspValIleProValArgArg 120
 |||||
 Db 1230 GGCCTCTCGACCTTTACCTGTGTACAGAGCAGCCGCGATGTCTATCCCGTGGCGCGCG 1289
 QY 121 GlyAspSerArgGlySerLeuLeuSerProAlaGlyHisAlaValGlyIlePheArgAlaIaValCys 160
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 Db 1290 GGTGTATGAGGAGGAGCGCTGCTGTGCGCCCGGCCCATTTCTCTACTTGAAGGCTCTCTCG 1349
 QY 141 GlyGlyProLeuLeuCysProAlaGlyHisAlaValGlyIlePheArgAlaIaValCys 160
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 Db 1350 GGGGTCCGCTGTGTGTCGCCCGGGGCCACCGCGTGGGCATATTTAGGCGCGCGGTGTGC 1409
 QY 161 ThrArgGlyValAlaIysAlaValAspPheIleProValGluSerLeuGluThrThrMet 180
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Db 1410 ACCCGTGGAGTGGCTAGAGGGGTGGACTTATCCCTGTGGAGAACCTAGAGACACCATG 1469
 QY 161 ArgSer 182
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 Db 1470 AGGTCC 1475

RESULT 7
 I09328
 LOCUS 109328
 DEFINITION Sequence 8 from Patent WO 8904669.
 ACCESSION 109328
 VERSION 109328.1 GI:587963
 KEYWORDS
 SOURCE Unknown.
 ORGANISM Unknown.
 REFERENCE 1 (bases 1 to 5360)
 AUTHORS Houghton,M., Choo,Q.-L. and Kuo,G.
 JOURNAL Patent: WO 8904669-A 8 01-JUN-1989;
 FEATURES
 location/Qualifiers
 1..5360
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 BASE COUNT 1061 a 1623 c 1533 g 1143 t
 ORIGIN

Alignment Scores:
 Pred. No.: 3,226-67 Length: 5360
 Score: 943.00 Matches: 179
 Percent Similarity: 100.00% Conservative: 3
 Best Local Similarity: 98.35% Mismatches: 0
 Query Match: 98.95% Indels: 0
 DB: 6 Gaps: 0

US-09-965-594-1 (1-182) x 109328 (1-5360)

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 |||||
 Db 990 AGCTTAACISGCCGAGACAAACACAGTGAAGGTGAGGTCCAGATTCGTACTGCT 1049
 QY 41 AlaGlnThrPheLeuAlaThrCysIleAsnGlyValCysTrpThrValCysHisGlyAla 60
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 Db 1050 GGCACAAACCTTCCTGCAACAGTGCATCAATGGGTGTGTGACTGTCTACCAAGGGGCC 1109
 QY 61 GlyThrArgThrIleAlaSerProLysGlyProValIleGlnMetTyrThrAsnValAsp 80
 |||||
 Db 1110 GGAACAGAGACCATCCGTCACCAAGGTCCTGTCTCATCCAGATGTATACCAATGTAGAC 1169
 QY 81 LysAspLeuValGlyTrpProAlaProGlnGlySerArgSerLeuThrProCysThrCys 100
 |||||
 Db 1170 CAAGACCTTGTGGGTGGGCCCTCCGCAAGGTAGCCGCTCATTCACACCTGCACTTGC 1229
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 |||||
 Db 1230 GGCCTCTCGACCTTTACCTGTGTACAGAGCAGCCGCGATGTCTATCCCGTGGCGCGCG 1289
 QY 121 GlyAspSerArgGlySerLeuLeuSerProAlaGlyHisAlaValGlyIlePheArgAlaIaValCys 160
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 Db 1290 GGTGTATGAGGAGGAGCGCTGCTGTGCGCCCGGCCCATTTCTCTACTTGAAGGCTCTCTCG 1349
 QY 141 GlyGlyProLeuLeuCysProAlaGlyHisAlaValGlyIlePheArgAlaIaValCys 160
 |||||
 Db 1350 GGGGTCCGCTGTGTGTCGCCCGGGGCCACCGCGTGGGCATATTTAGGCGCGCGGTGTGC 1409
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 |||||
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 QY 181 ArgSer 182
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Db 1470 AGGTCC 1475

RESULT 8
AR118692

LOCUS
DEFINITION Sequence 65 from patent US 6150087. linear PAT 16-MAY-2001
ACCESSION AR118692
VERSION AR118692.1 GI:14100602
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 6785)
AUTHORS Chien, D.Y.
TITLE NANV diagnostics and vaccines
JOURNAL Patent: US 6150087-A, 65 21-MAY-2000;
FEATURES
source location/Qualifiers
1..6785 /organism="unknown"
BASE COUNT 1392 a 2050 c 1914 g 1429 t
ORIGIN

Alignment Scores:
Pred. No.: 4.15e-67 Length: 6785
Score: 943.00 Matches: 179
Percent Similarity: 100.00% Conservative: 3
Best Local Similarity: 98.35% Mismatches: 0
Query Match: 98.95% Indels: 0
DB: 6 Gaps: 0

US-09-965-594-1 (1-192) x AR118692 (1-6785)

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QY 21 SerLeuThrGlyArgAspLysAsnGlnValGluGlyGluValGlnIleValSerThrAla 40
DB 1263 AGCCTAACTGGCGGGGACAAAACCAAGTGGAGGGTCCAGATTGTCTCAACTGCT 1322
QY 41 AlaGlnThrPheLeuAlaThrCysIleAsnGlyValCysTyrThrValTyrHisGlyAla 60
DB 1323 GCCCAAACTTCTGGCAACGTGCATCAATGGGGTGTCTGGACTGTCTACACGGGGCC 1382
QY 61 GlyThrArgThrIleAlaSerProLysGlyProValIleGlnMetTyrThrAsnValAsp 90
DB 1383 GGAACGAGGACCATCGCTACCCCAAGGCTCTGTATCCAGATGTAACCAATGTAGAC 1442
QY 81 LysAspLeuValGlyTyrProAlaProGlnGlySerArgSerLeuThrProCysThrCys 100
DB 1443 CAAGACCTTGTGGCTGGCGGCTCCGCAAGGTAGCGGCTCATTGACACCTGCACCTTC 1502
QY 101 GlySerSerAspLeuTyrLeuValThrArgHisAlaAspValIleProValArgArgArg 120
DB 1503 GGCTCTCGGACCTTACCTGGTCCAGGACCGCGCATGTCATTCGCGCGCGCGG 1562
QY 121 GlyAspSerArgGlySerLeuLeuSerProArgProIleSerTyrLeuLysGlySerSer 140
DB 1563 GGTGATAGAGGGGACGCTGCTGCCCGGCGCCCAATTTCTTACTTGAAGGCTCCTCG 1622
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QY 161 ThrArgGlyValAlaLysAlaValAspPheIleProValGluSerLeuGluThrThrMet 180
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QY 181 ArgSer 182
DB 1743 AGGTCC 1748

RESULT 10
109329

LOCUS
DEFINITION Sequence 10 from Patent WO 8904669. linear PAT 02-DEC-1994

106440

LOCUS
DEFINITION Sequence 54 from Patent EP 0318216. linear PAT 02-DEC-1994
ACCESSION 106440
VERSION 106440.1 GI:590312
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 6785)
AUTHORS Houghton, M., Choo, Q.-L. and Kuo, G.
TITLE NANV diagnostics and vaccines
JOURNAL Patent: EP 0318216-A1 54 31-MAY-1989;
FEATURES
source location/Qualifiers
1..6785 /organism="unknown"
BASE COUNT 1392 a 2050 c 1914 g 1429 t
ORIGIN

Alignment Scores:
Pred. No.: 4.15e-67 Length: 6785
Score: 943.00 Matches: 179
Percent Similarity: 100.00% Conservative: 3
Best Local Similarity: 98.35% Mismatches: 0
Query Match: 98.95% Indels: 0
DB: 6 Gaps: 0

US-09-965-594-1 (1-182) x 106440 (1-6785)

QY 1 MetAlaProIleThrAlaTyrAlaGlnGlnThrArgGlyLeuLeuGlyCysIleIleThr 20
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DB 1323 GCCCAAACTTCTGGCAACGTGCATCAATGGGGTGTCTGGACTGTCTACACGGGGCC 1382
QY 61 GlyThrArgThrIleAlaSerProLysGlyProValIleGlnMetTyrThrAsnValAsp 80
DB 1383 GGAACGAGGACCATCGCTACCCCAAGGCTCTGTATCCAGATGTAACCAATGTAGAC 1442
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DB 1443 CAAGACCTTGTGGCTGGCGGCTCCGCAAGGTAGCGGCTCATTGACACCTGCACCTTC 1502
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DB 1563 GGTGATAGAGGGGACGCTGCTGCCCGGCGCCCAATTTCTTACTTGAAGGCTCCTCG 1622
QY 141 GlyGlyProLeuLeuCysProAlaGlyHisAlaValGlyIlePheArgAlaAlaValCys 160
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DB 1743 AGGTCC 1748

RESULT 10
109329

LOCUS
DEFINITION Sequence 10 from Patent WO 8904669. linear PAT 02-DEC-1994

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ACCESSION      I09329
VERSION        I09329.1  GI:587964
KEYWORDS       (unknown)
SOURCE         Unclassified.
ORGANISM       Unclassified.
REFERENCE      1 (bases 1 to 6785)
AUTHORS        Broughton, M., Choo, Q.-K., and Kuo, G.
JOURNAL        Patent: WO 8904689-A 10.01-JUN-1989;
FEATURES       Location/Qualifiers
               source          1..6785
               /organism="unknown"
BASE COUNT     1392 a 2050 c 1914 g 1423 t
ORIGIN
Alignment Scores:
Pred. No.:      4,156-67      Length:      6785
Score:          943.00      Matches:    179
Percent Similarity: 100.00%      Conservatives: 3
Best Local Similarity: 98.95%      Mismatches: 0
Query Match:    98.95%      Indels:    0
DB:              6      Gaps:      0
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DB      1263 AGCCTAATCGCGGACAAAACCAACCAAGTGGAGGTGAGTCCAGATTGTGTCAACTGCT 1322
QY      41 AlaGlnThrPheLeuAlaThrCysIleAsnGlyValCysTrpThrValTyrHisGlyAla 60
DB      1323 GCCCAAACTCTCTGCAACGTCATCAATAGGAGGTGTGTGCACTGTCTTACCAAGGGGCC 1382
QY      61 GlyThrArgThrIleAlaSerProLysGlyProValIleLeuMetTyrThrAsnValAsp 80
DB      1393 GGAACAGAGACCATCCGTCACCAAGGTCCTGTGTCATCCAGATGATACCAATGTAGAC 1442
QY      81 LysAspLeuValClyTrpProAlaProGlnGlySerArgSerSerLeuThrProCysThrCys 100
DB      1443 CAAGACCTTGTGGGTGGCGCGCTCGGCAAGGATGAGCGGTCTATGACACCTGGCACTTGC 1502
QY      121 GlySerSerAspLeuThrLeuValThrArgHisAlaAspValIleProValArgArgArg 120
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QY      121 GlyAspSerArgGlySerLeuLeuSerProArgProIleSerTyrLeuLysGlySerSer 140
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QY      141 GlyGlyProLeuLeuCysProAlaGlyHisAlaValGlyIlePheArgAlaAlaValCys 160
DB      1623 GGGGGTCCGCTGTGTGGCGCGCGGCGGACAGCGCGCGGCGGATATTTAGGCGCGCGGTGTC 1682
QY      161 ThrArgGlyValAlaLysAlaValAspPheIleProValGluSerLeuGluThrThrMet 180
DB      1683 ACCCGTGGAGTGGCTAAGCGGTGGGACTTATCTCGGTGTGAGAACCTAGAGATAACCAAG 1742
QY      181 ArgSer 182
DB      1743 AGGTCC 1748
RESULT 11
LOCUS         AR118696
DEFINITION    Sequence 74 from patent US 6150087.
ACCESSION     AR118696
VERSION       AR118696.1  GI:14106606
KEYWORDS      (unknown)
SOURCE        Unclassified.
REFERENCE     1 (bases 1 to 7310)
LOCUS         T09331
DEFINITION    Sequence 15 from Patent WO 8904669.
ACCESSION     T09331
VERSION       T09331.1  GI:587966
KEYWORDS      (unknown)
SOURCE        Unclassified.
REFERENCE     1 (bases 1 to 7310)
ORGANISM      Unknown.

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Db 1786 AGCCCTAACCTGGCGGCGCAAAACCAAGTGGAGGCTGAGGTCCAGATATGTCACACTGCT 1847
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QY 61 GlyThrArgThrIleAlaSerProLysGlyProValIleGlnMetTyrThrAsnValAsp 80
Db 1908 GGAACAGCAGCACCCTCGCTACCAAGGCTCTGTATCCAGATGATACCAATATGAGAC 1967
QY 81 LysAspLeuValGlyTrpProAlaProGlnGlySerArgSerLeuThrProCysThrCys 100
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QY 161 ThrArgGlyValAlaAlaLysAlaValAspPheIleProValGluSerLeuGluThrThrMet 180
Db 2208 ACCGTGTGAGTGGCTAAGCGGTGGACTTTATCCCTGTGGAGACCTAGAGACACCAATG 2267
QY 181 ArgSer 182
Db 2268 AGGTCC 2273
RESULT 14
AR118703
LOCUS AR118703 8316 bp DNA linear PAT 16-MAY-2001
DEFINITION Sequence 88 from patent US 6150087.
ACCESSION AR118703
VERSION AR118703.1 GI:14100612
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 8316)
AUTHORS Chien,D.Y.
TITLE NANBV diagnostics and vaccines
JOURNAL Patent: US 6150087-A 88 21-NOV-2000;
FEATURES
    location/Qualifiers
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BASE COUNT 1671 a 2529 c 2345 g 1771 t
ORIGIN
Alignment Scores:
Pred. No.: 5,17e-67 Length: 8316
Score: 943.00 Matches: 179
Percent Similarity: 100.00% Conservative: 3
Best Local Similarity: 98.35% Mismatches: 0
Query Match: 98.95% Indels: 0
DB: 6 Gaps: 0
US-09-965-594-1 (1-182) x AR118703 (1-8316)
QY 1 MetAlaProIleThrAlaTyrAlaGlnGlnThrArgGlyLeuLeuGlyCysIleIleThr 20
Db 2734 CTGGCGCCCATCAGCGGTAGCGCCAGCAGCAAGGCGGCTCTAGGTCATATACACC 2793
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QY 21 SerLeuThrGlyArgAspLysAsnGlyValGluGlyGluValGlnIleValSerThrAla 40
Db 2794 AGCCCTAACCTGGCGGCGCAAAACCAAGTGGAGGCTGAGGTCCAGATATGTCACACTGCT 2853
QY 41 AlaGlnThrPheLeuAlaThrCysIleAsnGlyValCysTrpThrValTyrHisGlyAla 60
Db 2854 GCCCAAACTTCTCGGCAACGTGATCAATGGGCTGCTGACTGTCTACACAGCGGCGCC 2913
QY 61 GlyThrArgThrIleAlaSerProLysGlyProValIleGlnMetTyrThrAsnValAsp 80
Db 2914 GGAACAGCAGCACCCTCGCTACCAAGGCTCTGTATCCAGATGATACCAATATGAGAC 2973
QY 81 LysAspLeuValGlyTrpProAlaProGlnGlySerArgSerLeuThrProCysThrCys 100
Db 2974 CAAGACCTTGTGGCTGGCGCGCTCGGCAAGGTAGCGGCTCATACACCCCTGCACTTGC 3033
QY 101 GlySerSerAspLeuTyrLeuValThrArgHisAlaAspValIleProValArgArgArg 120
Db 3034 GGCCTCTCGGACCTTACCTGTGTACAGCAGCAGCGCGATGTCTATCCCTGGCGCGCGG 3093
QY 121 GlyAspSerArgGlySerLeuLeuSerProAlaProLysSerTyrLeuLysGlySer 140
Db 3094 GGTGATAGCAGGGCAGCGTGTGTGGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCG 3153
QY 141 GlyGlyProLeuLeuCysProAlaGlyHisAlaValGlyIlePheArgAlaAlaValCys 160
Db 3154 GGGGTTCGCTGTGTGGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCG 3213
QY 161 ThrArgGlyValAlaAlaLysAlaValAspPheIleProValGluSerLeuGluThrThrMet 180
Db 3214 ACCGTGTGAGTGGCTAAGCGGTGGACTTTATCCCTGTGGAGACCTAGAGACACCAATG 3273
QY 181 ArgSer 182
Db 3274 AGGTCC 3279
RESULT 15
AR118728
LOCUS AR118728 8987 bp DNA linear PAT 16-MAY-2001
DEFINITION Sequence 137 from patent US 6150087.
ACCESSION AR118728
VERSION AR118728.1 GI:14100618
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 8987)
AUTHORS Chien,D.Y.
TITLE NANBV diagnostics and vaccines
JOURNAL Patent: US 6150087-A 137 21-NOV-2000;
FEATURES
    location/Qualifiers
        source
            1..8987
            /organism="unknown"
BASE COUNT 1807 a 2735 c 2547 g 1898 t
ORIGIN
Alignment Scores:
Pred. No.: 5,62e-67 Length: 8987
Score: 943.00 Matches: 179
Percent Similarity: 100.00% Conservative: 3
Best Local Similarity: 98.35% Mismatches: 0
Query Match: 98.95% Indels: 0
DB: 6 Gaps: 0
US-09-965-594-1 (1-182) x AR118728 (1-8987)
QY 1 MetAlaProIleThrAlaTyrAlaGlnGlnThrArgGlyLeuLeuGlyCysIleIleThr 20
Db 3076 CTGGCGCCCATCAGCGGTAGCGCCAGCAGCAAGGCGGCTCTAGGTCATATACACC 3135
QY 21 SerLeuThrGlyArgAspLysAsnGlnValGluGlyGluValGlnIleValSerThrAla 40
Db 3136 AGCCTAACCTGGCGGCGCAAAACCAAGTGGAGGCTGAGGTCCAGATATGTCACACTGCT 3195
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```
QY 41 AlaGlnThrPheLeuAlaThrCysIleAsnGlyValCysTrpThrValTyrHisGlyAla 60
Db 3196 GCCCAACCTTCCCTGGCAACATGCATCAATGGGTGGCTGGACTGCTACCCAGGGGAC 3255
QY 61 GlyThrArgThrIleAlaSerProLysGlyProValIleGlnMetTyrThrAsnValAsp 80
Db 3256 GGAACGAGGACCATCGGTCAACCAAGGGTCTCTGTCATCCAGATGATACCAATCTAGAC 3315
QY 81 LysAspLeuValGlyTyrProAlaProGlnGlySerArgSerLeuThrProCysThrCys 100
Db 3316 CAAGACCTTGGGCTCGCCCGCTCCGCAAGTAGCCGCTAATTSACACCTGCACCTGC 3375
QY 101 GlySerSerAspLeuTyrLeuValThrArgHisAlaAspValIleProValArgArgArg 120
Db 3376 GGCTCTCGGACCTTACCTGGTACAGAGACAGCGCATGTCTATCCGCTGGCGCGCGCG 3435
QY 121 GlyAspSerArgGlySerLeuLeuSerProArgProIleSerTyrLeuGlyLysGlySerSer 140
Db 3436 GGTGATACGAGGGACGCTACTGTGCGCCGCGGCGCATTTCTACTTGAAAGGCTCTGG 3495
QY 141 GlyGlyProLeuLeuCysProAlaGlyHisAlaValGlyIlePheAsnAlaAlaValCys 160
Db 3496 GGGGSHYGGCTGTGTGTCGCGCGGCGACGCTGTGSGCATTTTAGGSCVARGSTETGG 3555
QY 161 ThrArgGlyValAlaLysAlaValAspPheIleProValIleIleSerLeuS-uthrThrMet 180
Db 3556 ACCGTGAGTGGCTAAGCGGTGSACTTTATCCCTGTGAGAGACCTAGAGACACCATG 3615
QY 181 ArgSer 182
Db 3616 AGGTCC 3621
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Search completed: September 27, 2003, 13:32:26
Job time : 3361 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2003 Computer Ltd.

OM protein - protein search, using sw model

Run on: September 27, 2003, 11:35:53 : Search time 107 Seconds
(without alignments)
436,931 Million cell updates/sec

Title: US-09-965-594-1

Perfect score: 953

Sequence: 1 MAPITAYAQTRGLGCGIT.....GVAXAVDFIPVESLEIMKPS 182

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 830525 seqs, 258052604 residues

Total number of hits satisfying chosen parameters: 830525

Minimum DB seq length: 6

Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%

Maximum Match 99%

Listing first 45 summaries

Database :

SPTREMBL_23.*

- 1: sp_archaea.*
- 2: sp_bacteria.*
- 3: sp_fungi.*
- 4: sp_human.*
- 5: sp_invertebrate.*
- 6: sp_mammal.*
- 7: sp_mbc.*
- 8: sp_oragaele.*
- 9: sp_phage.*
- 10: sp_plant.*
- 11: sp_rodent.*
- 12: sp_virus.*
- 13: sp_vertebrate.*
- 14: sp_unclassified.*
- 15: sp_virus.*
- 16: sp_bacteriap.*
- 17: sp_archaeap.*

Pred. No. is the number of results predicted by glance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match %	Length	DB ID	Description
1	945	99.2	181	12 Q91RS8	Q91RS8 hepatitis C
2	945	99.2	181	12 Q91RT5	Q91RT5 hepatitis C
3	943	99.0	2436	12 Q81758	Q81758 hepatitis C
4	943	99.0	3011	12 Q91FES	Q91FES hepatitis C
5	942	98.8	181	12 Q91RS3	Q91RS3 hepatitis C
6	942	98.8	181	12 Q91RS1	Q91RS1 hepatitis C
7	942	98.8	181	12 Q91RQ8	Q91RQ8 hepatitis C
8	942	98.8	181	12 Q91RT1	Q91RT1 hepatitis C
9	940	98.6	181	12 Q91RS6	Q91RS6 hepatitis C
10	940	98.6	181	12 Q91RS9	Q91RS9 hepatitis C
11	939	98.5	181	12 Q91RS3	Q91RS3 hepatitis C
12	939	98.5	3011	12 Q93463	Q93463 hepatitis C
13	938	98.4	181	12 Q91RT4	Q91RT4 hepatitis C
14	938	98.4	181	12 Q91RS8	Q91RS8 hepatitis C
15	938	98.4	181	12 Q91RT3	Q91RT3 hepatitis C
16	938	98.4	181	12 Q91RS5	Q91RS5 hepatitis C

17	936	98.4	181	12 Q91RS7	Q91RS7 hepatitis C
18	936	98.4	181	12 Q91RT0	Q91RT0 hepatitis C
19	937	98.3	3021	12 Q36579	Q36579 hepatitis C
20	936	98.2	181	12 Q91RS5	Q91RS5 hepatitis C
21	936	98.2	181	12 Q91RS4	Q91RS4 hepatitis C
22	935	98.1	181	12 Q91RT6	Q91RT6 hepatitis C
23	934	98.0	181	12 Q91RT9	Q91RT9 hepatitis C
24	933	97.9	181	12 Q91RS4	Q91RS4 hepatitis C
25	933	97.9	181	12 Q91RS9	Q91RS9 hepatitis C
26	933	97.9	181	12 Q91RS0	Q91RS0 hepatitis C
27	933	97.9	3011	12 Q91RS9	Q91RS9 hepatitis C
28	932	97.8	181	12 Q91RS2	Q91RS2 hepatitis C
29	931	97.7	181	12 Q91RR7	Q91RR7 hepatitis C
30	931	97.7	3011	12 Q91RT6	Q91RT6 hepatitis C
31	930	97.6	181	12 Q91RT2	Q91RT2 hepatitis C
32	930	97.6	181	12 Q91RS1	Q91RS1 hepatitis C
33	930	97.6	181	12 Q91RS2	Q91RS2 hepatitis C
34	930	97.6	181	12 Q91RS2	Q91RS2 hepatitis C
35	930	97.6	3011	12 Q36608	Q36608 hepatitis C
36	930	97.6	3015	12 Q9PWK5	Q9PWK5 hepatitis C
37	930	97.6	3015	12 Q9PWJ9	Q9PWJ9 hepatitis C
38	928	97.4	181	12 Q91RS6	Q91RS6 hepatitis C
39	927	97.3	181	12 Q91RT7	Q91RT7 hepatitis C
40	925	97.1	181	12 Q91RS0	Q91RS0 hepatitis C
41	925	97.1	181	12 Q91RT8	Q91RT8 hepatitis C
42	924	97.0	3011	12 Q36609	Q36609 hepatitis C
43	919	96.4	3011	12 Q36610	Q36610 hepatitis C
44	909	95.4	3011	12 Q81754	Q81754 hepatitis C
45	903	94.8	3010	12 Q9J3G9	Q9J3G9 hepatitis C

ALIGNMENTS

RESULT 1

Q91RS8
ID Q91RS8 PRELIMINARY: PRT: 181 AA.

AC Q91RS8: 01-DEC-2001 (TRENBLrel. 19, Created)

DT 01-DEC-2001 (TRENBLrel. 19, Last sequence update)

DT 01-MAR-2003 (TRENBLrel. 23, Last annotation update)

DE NS3 protease (Fragment)

OS Hepatitis C virus.

OC Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;

CC Hepacivirus.

GN NCBI_TaxID:11103;

RN 11

RF SEQUENCE FROM N.A.

RC STRAIN-Pt.1Y;

KA Holland-Staley C.A., Kovari L.C., Golenberg E., Mayers D.L.;

RT "Genetic Diversity and response to IFN of the NS3 Protease Gene from

RI Clinical Strains of the Hepatitis C Virus."

RL Submitted (APR-2001) to the EMBL/GenBank/DBJ databases.

DR EMBL; AF369235; AAK34560.1; -

DR InterPro; IPR004109; HCV_NS3.

DR Pfam; PFC2907; HCV_NS3; 1.

KW Protease.

FI NON_TER 1 1

FT NON_TER 181 181

SQ SEQUENCE 181 AA: 19130 KW: 85D91869299B7C35 CRC64:

Query Match 99.2%; Score 945; DB 12: Length 181;

Best Local Similarity 99.4%; Pred No. 1,9e-88;

Matches 180; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 2 APITAYAQTRGLGCGITSLTGRDKNOVEQIVSTAAQTFLAICINGCVTVYHGAG 61
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DB 1 APITAYAQTRGLGCGITSLTGRDKNOVEQIVSTAAQTFLAICINGCVTVYHGAG 60
QY 62 TTTIASPGVPIQMTYNVDKDLGVNPAQGSRSLSLTPCTCGSSDLYLVTRHADVIPVRRG 121
|||||
DB 61 TTTIASPGVPIQMTYNVDKDLGVNPAQGSRSLSLTPCTCGSSDLYLVTRHADVIPVRRG 120

QY 122 DSRGSLSPRPISVYKSGSGGGLCPAGHAGVGFRAAVCTRGVAKAVDFIPVESLETTMR 181
 DB DSRGSLSPRPISVYKSGSGGGLCPAGHAGVGFRAAVCTRGVAKAVDFIPVESLETTMR 180
 QY 182 S 182
 DB 181 S 181

RESULT 2

Q91RT5 ID Q91RT5 PRELIMINARY; PRI: 181 AA.
 AC Q91RT5;
 DT 01-DEC-2001 (Tremblrel. 19, Created)
 DT 01-DEC-2001 (Tremblrel. 19, Last sequence update)
 DT 01-MAR-2003 (Tremblrel. 23, Last annotation update)
 DE NS3 protease (Fragment).
 OS Hepatitis C virus.
 OC Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
 OC Hepacivirus.
 OX NCBI_TaxID=11103;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN:P-4;
 RA Holland-Staley C.A., Kovari L.C., Giesberg E., Meyers D.L.;
 RT "Genetic Diversity and Response to IFN of the NS3 Protease Gene from
 RT Clinical Strains of the Hepatitis C Virus."
 RL Submitted (Apr-2001) to the EMBL/GenBank/DDBJ databases.
 DR EMBL: AF169218; AA54543.1;
 DR InterPro: IPR004139; HCV_NS3.
 DR Pfam: PF02907; HCV_NS3; 1.
 KW Protease.
 FT NON_TER 181 181
 FT NON_TER 181 181
 SQ SEQUENCE 181 AA: 19:30 MW: 85D9:564294B3C35 CRC64:

Query Match 99.28; Score 9457; DR 12; Length 182;
 Best Local Similarity 99.44; Pred. No. 1.9e-88;
 Matches 180; Conservative 1; Mismatches 0; Indels 0; Gaps 0;
 QY 2 APIITAYAOQTGGLGCIITSLTGDRKNQVEGEVQIVSTAAQTFLATCINGVCTVYHGA 61
 DB 1 APIITAYAOQTGGLGCIITSLTGDRKNQVEGEVQIVSTAAQTFLATCINGVCTVYHGA 60
 QY 62 TRIASPKGPIQVIQYINVDKDLVGMWPAQSGRSITPCTCGSSDLYLVTRHADVIPVRR 121
 DB 61 TRIASPKGPIQVIQYINVDKDLVGMWPAQSGRSITPCTCGSSDLYLVTRHADVIPVRR 120
 QY 122 DSRGSLSPRPISVYKSGSGGGLCPAGHAGVGFRAAVCTRGVAKAVDFIPVESLETTMR 181
 DB 121 DSRGSLSPRPISVYKSGSGGGLCPAGHAGVGFRAAVCTRGVAKAVDFIPVESLETTMR 180
 QY 182 S 182
 DB 181 S 181

RESULT 3

Q81756 ID Q81756 PRELIMINARY; PRI: 2436 AA.
 AC Q81756;
 DT 01-NOV-1996 (Tremblrel. 01, Created)
 DT 01-NOV-1996 (Tremblrel. 01, Last sequence update)
 DT 01-MAR-2003 (Tremblrel. 23, Last annotation update)
 DE Genome polyprotein (Fragment).
 OS Hepatitis C virus.
 OC Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
 OC Hepacivirus.
 OX NCBI_TaxID=11103;
 RN [1]
 RP SEQUENCE FROM N.A.
 RA Choo Q.-L., Richman K., Han J.;
 RT "The nucleotide sequence of the Hepatitis C viral genome."

RL Submitted (May-1990) to the EMBL/GenBank/DDBJ databases.
 DR EMBL: M32084; AAA45677.1;
 DR HSP: P27958; IAIV
 DR InterPro: IPR001410; DEAD.
 DR InterPro: IPR002531; HCV_NS1.
 DR InterPro: IPR002518; HCV_NS2.
 DR InterPro: IPR004109; HCV_NS3.
 DR InterPro: IPR000745; HCV_NS4a.
 DR InterPro: IPR001490; HCV_NS4b.
 DR InterPro: IPR002868; HCV_NS5a.
 DR InterPro: IPR002166; HCV_RDRP.
 DR InterPro: IPR001650; Helicase_C.
 DR InterPro: IPR007095; RNA_po_DS_PS.
 DR InterPro: IPR007094; RNA_po_PS_vir.
 DR Pfam: PF01560; HCV_NS1; 1.
 DR Pfam: PF01518; HCV_NS2; 1.
 DR Pfam: PF02907; HCV_NS3; 1.
 DR Pfam: PF01006; HCV_NS4a; 1.
 DR Pfam: PF01001; HCV_NS4b; 1.
 DR Pfam: PF01506; HCV_NS5a; 1.
 DR Pfam: PF00271; Helicase_C; 1.
 DR Pfam: PF00598; Viral_RDRP; 1.
 DR Pfam: PF01866; HCV_NS1; 1.
 DR ProDom: PD186062; HCV_NS1; 1.
 DR SMART: SM03487; REXDC; 1.
 DR PROSITE: PS00507; RDRP_POSITIVE; 1.
 DR PROSITE: PS0521; RDRP_VIRAL; 1.
 KW ATP-binding; Coat protein; Envelope protein; Glycoprotein; Helicase;
 KW Hydrolase; Nonstructural protein; Polyprotein;
 KW RNA-directed RNA polymerase; Transferase; Transmembrane.
 FT NON_TER 2436 2436
 FT NON_TER 2436 2436
 SQ SEQUENCE 2436 AA: 264734 MW: D79872900BE3125 CRC64:

Query Match 99.04; Score 943; DR 12; Length 2436;
 Best Local Similarity 98.48; Pred. No. 7.1e-87;
 Matches 279; Conservative 3; Mismatches 0; Indels 0; Gaps 0;
 QY 1 MAPITAYAOQTGGLGCIITSLTGDRKNQVEGEVQIVSTAAQTFLATCINGVCTVYHGA 60
 DB 576 LAPITAYAOQTGGLGCIITSLTGDRKNQVEGEVQIVSTAAQTFLATCINGVCTVYHGA 635
 QY 61 GTETASPKGPIQVIQYINVDKDLVGMWPAQSGRSITPCTCGSSDLYLVTRHADVIPVRR 120
 DB 636 GTETASPKGPIQVIQYINVDKDLVGMWPAQSGRSITPCTCGSSDLYLVTRHADVIPVRR 695
 QY 121 GDSRGSLLSPRPISVYKSGSGGGLCPAGHAGVGFRAAVCTRGVAKAVDFIPVESLETTM 180
 DB 696 GDSRGSLLSPRPISVYKSGSGGGLCPAGHAGVGFRAAVCTRGVAKAVDFIPVESLETTM 755
 QY 181 RS 182
 DB 756 RS 757

RESULT 4

Q91FE5 ID Q91FE5 PRELIMINARY; PRI: 3011 AA.
 AC Q91FE5;
 DT 01-OCT-2000 (Tremblrel. 15, Created)
 DT 01-OCT-2000 (Tremblrel. 15, Last sequence update)
 DT 01-MAR-2003 (Tremblrel. 23, Last annotation update)
 DE Genome polyprotein.
 OS Hepatitis C virus.
 OC Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
 OC Hepacivirus.
 OX NCBI_TaxID=11103;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=21262212; PubMed=11369872;
 RA Lanford K.E., Lee H., Chavez D., Guerra B., Brasky K.M.;
 RT "Infectious cDNA clone of the hepatitis C virus genotype 1 prototype
 sequence."
 RL J. Gen. Virol. 82:1291-1297(2001).


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Db 61 TTTIATSPKGPVQIOMYTNVQDLVGVWPAPQAGRSLLTCTCGSSDLVLTTRHADVIPVRRG 120
QY 122 DSRGSLSPRPISYLGSSGGPLLCAGHAGVFRAAVCTRGVAKAVDFIPVESLETTMR 181
DB 121 DSRGSLSPRPISYLGSSGGPLLCAGHAGVFRAAVCTRGVAKAVDFIPVESLETTMR 180
QY 182 S 182
DB 181 S 181

RESULT 10
Q91RS9 PRELIMINARY: PRT: 181 AA.
AC Q91RS9;
DT 01-DEC-2001 (TrEMBLrel. 19, Created)
DT 01-DEC-2001 (TrEMBLrel. 19, Last sequence update)
DT 01-MAR-2003 (TrEMBLrel. 23, Last annotation update)
DE NS3 protease (Fragment).
OS Hepatitis C virus.
OC Hepacivirus.
OC Viruses; ssRNA positive-strand viruses, no DNA stage: Flaviviridae;
OX NCBI_TaxID=11103;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=Pt.174;
RA Holland-Staley C.A., Kovari L.C., Golenberg E., Mayers D.L.;
RT "Genetic Diversity and response to IFN of the NS3 Protease Gene from
Clinical Strains of the Hepatitis C Virus.";
RL Submitted (APR-2001) to the EMBL/GenBank/DBSJ databases.
DR EMBL: AF369224; AAK54549.1;
DR InterPro: IPR004109; HCV_NS3.
DR Pfam: PF02907; HCV_NS3; 1.
KW Protease.
FT NON_TER 1 181
FT NON_TER 181 S 181
SQ SEQUENCE 181 AA: 19131 MW: 89076279 DBD635 CRC64;

Query Match 98.5%; Score 940; DB 12; Length 181;
Best Local Similarity 98.9%; Pred. No. 6e-88;
Matches 179; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

QY 2 APITAYAOQTGRLGCGITISLTGRKKNQVEGVQIVSTAAQTFLATCINGVCWTVYHGAG 61
DB 1 APITAYAOQTGRLGCGITISLTGRKKNQVEGVQIVSTAAQTFLATCINGVCWTVYHGAG 60
QY 62 TTTIATSPKGPVQIOMYTNVQDLVGVWPAPQAGRSLLTCTCGSSDLVLTTRHADVIPVRRG 121
DB 61 TTTIATSPKGPVQIOMYTNVQDLVGVWPAPQAGRSLLTCTCGSSDLVLTTRHADVIPVRRG 120
QY 122 DSRGSLSPRPISYLGSSGGPLLCAGHAGVFRAAVCTRGVAKAVDFIPVESLETTMR 181
DB 121 DSRGSLSPRPISYLGSSGGPLLCAGHAGVFRAAVCTRGVAKAVDFIPVESLETTMR 180
QY 182 S 182
DB 181 S 181

RESULT 11
Q91RS3 PRELIMINARY: PRT: 181 AA.
AC Q91RS3;
DT 01-DEC-2001 (TrEMBLrel. 19, Created)
DT 01-DEC-2001 (TrEMBLrel. 19, Last sequence update)
DT 01-MAR-2003 (TrEMBLrel. 23, Last annotation update)
DE NS3 protease (Fragment).
OS Hepatitis C virus.
OC Viruses; ssRNA positive-strand viruses, no DNA stage: Flaviviridae;
OX NCBI_TaxID=11103;
RN [1]
RP SEQUENCE FROM N.A.
```

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RC STRAIN=Pt.24;
RA Holland-Staley C.A., Kovari L.C., Golenberg E., Mayers D.L.;
RT "Genetic Diversity and response to IFN of the NS3 Protease Gene from
Clinical Strains of the Hepatitis C Virus.";
RL Submitted (APR-2001) to the EMBL/GenBank/DBSJ databases.
DR EMBL: AF369224; AAK54555.1;
DR InterPro: IPR004109; HCV_NS3.
DR Pfam: PF02907; HCV_NS3; 1.
KW Protease.
FT NON_TER 1 181
FT NON_TER 181 S 181
SQ SEQUENCE 181 AA: 19132 MW: 890805F3A395250 CRC64;

Query Match 98.5%; Score 939; DB 12; Length 181;
Best Local Similarity 98.3%; Pred. No. 7.6e-88;
Matches 178; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 2 APITAYAOQTGRLGCGITISLTGRKKNQVEGVQIVSTAAQTFLATCINGVCWTVYHGAG 61
DB 1 APITAYAOQTGRLGCGITISLTGRKKNQVEGVQIVSTAAQTFLATCINGVCWTVYHGAG 60
QY 62 TTTIATSPKGPVQIOMYTNVQDLVGVWPAPQAGRSLLTCTCGSSDLVLTTRHADVIPVRRG 121
DB 61 TTTIATSPKGPVQIOMYTNVQDLVGVWPAPQAGRSLLTCTCGSSDLVLTTRHADVIPVRRG 120
QY 122 DSRGSLSPRPISYLGSSGGPLLCAGHAGVFRAAVCTRGVAKAVDFIPVESLETTMR 181
DB 121 DSRGSLSPRPISYLGSSGGPLLCAGHAGVFRAAVCTRGVAKAVDFIPVESLETTMR 180
QY 182 S 182
DB 181 S 181

RESULT 12
Q93463 PRELIMINARY: PRT: 3011 AA.
AC Q93463;
DT 01-NOV-1996 (TrEMBLrel. 01, Created)
DT 01-NOV-1996 (TrEMBLrel. 01, Last sequence update)
DT 01-MAR-2003 (TrEMBLrel. 23, Last annotation update)
DE Genome polyprotein.
OS Hepatitis C virus.
OC Viruses; ssRNA positive-strand viruses, no DNA stage: Flaviviridae;
OX NCBI_TaxID=11103;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=HC-J1;
RX MEDLINE-91013116; PubMed-2170712;
RA Okamoto H., Okada S., Sugiyama Y., Yotsuroto S., Tanaka T.,
RA Yoshizawa H.;
RC "The 5'-terminal sequence of the hepatitis C virus genome.";
RC Jpn. J. Exp. Med. 60:167-177(1990).
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=HC-J1;
RX MEDLINE-92044440; PubMed-1658196;
RA Okamoto H., Okada S., Sugiyama Y., Kurai K., Iizuka H., Machida A.,
RA Miyakawa Y., Mayumi M.;
RT "Nucleotide sequences of the genomic RNA of hepatitis C virus isolated
from a human carrier: comparison with reported isolates for conserved
and divergent regions.";
RC J. Gen. Virol. 72:2697-2704(1991).
RN [3]
RP SEQUENCE FROM N.A.
RX MEDLINE-93117129; PubMed-1335573;
RA Okamoto H., Kanai N., Mishihiro S.;
RT "Full-length nucleotide sequence of a Japanese hepatitis C virus
isolate (HC-J1) with high homology to USA isolates.";
RL Nucleic Acids Res. 20:6410-6410(1992).
RN [4]
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RP SEQUENCE FROM N.A.
RC STRAIN=HC-J1;
RA Okamoto H.;
RL Submitted (DEC-1992) to the EMBL/GenBank/DBJ databases.
RN [5]
RP SEQUENCE FROM N.A.
RC STRAIN=HC-J1;
RX MEDLINE=94174722; PubMed=7510436;
RA Mink M., Benichou S., Madaule P., Hollais P., Prince A.,
RA Inchausti G.;
RI "Characterization and mapping of a B-cell immunogenic domain in
RT hepatitis C virus E2 glycoprotein using a yeast peptide library.";
RL Virology 200;246-255(1994).
CC !- SUBUNIT: THE VIRION OF THIS VIRUS IS A NUCLEOCAPSID COVERED BY A
CC LIPOPROTEIN ENVELOPE. THE ENVELOPE CONSISTS OF TWO PROTEINS: A
CC PROTEIN M AND GLYCOPROTEIN E. THE NUCLEOCAPSID IS A COMPLEX OF
CC PROTEIN C AND MRNA (BY SIMILARITY).
CC EMBL: D10749; BAA01582.1; -.
DR HSP: P27958; 1HE1.
DR InterPro: IPR001410; DEAD.
DR InterPro: IPR002522; HCV_capsid.
DR InterPro: IPR002521; HCV_core.
DR InterPro: IPR002519; HCV_env.
DR InterPro: IPR002531; HCV_NS2.
DR InterPro: IPR002518; HCV_NS2.
DR InterPro: IPR004109; HCV_NS3.
DR InterPro: IPR000745; HCV_NS4a.
DR InterPro: IPR001490; HCV_NS4b.
DR InterPro: IPR002868; HCV_NS5a.
DR InterPro: IPR002166; HCV_RDRP.
DR InterPro: IPR001650; Helicase_C.
DR InterPro: IPR007095; RNA_pol_DS_Ps.
DR InterPro: IPR007094; RNA_pol_PSVir.
DR Pfam: PF01543; HCV_capsid; 1.
DR Pfam: PF01542; HCV_core; 1.
DR Pfam: PF01539; HCV_env; 1.
DR Pfam: PF01560; HCV_NS1; 1.
DR Pfam: PF01538; HCV_NS2; 1.
DR Pfam: PF02907; HCV_NS3; 1.
DR Pfam: PF01005; HCV_NS4a; 1.
DR Pfam: PF01001; HCV_NS4b; 1.
DR Pfam: PF01506; HCV_NS5a; 1.
DR Pfam: PF00271; Helicase_C; 1.
DR Pfam: PF00598; Viral_Rdrp; 1.
DR ProDom: PD186062; HCV_NS1; 1.
DR SMART: SM00487; DEXDC; 1.
DR PROSITE: PS00407; RDRP_POSITIVE; 1.
DR PROSITE: PS0521; RDRP_VIRAL; 1.
KW ATP-binding; Coat protein; Envelope protein; Glycoprotein; Helicase;
KW Hydrolase; Nonstructural protein; Polyprotein;
KW RNA-directed RNA polymerase; Transferase; Transmembrane.
SQ SEQUENCE 3011 AA; 327112 MW; 97E9052C02504535 CRC64;

Query Match 98.5%; Score 930; DB 12; Length 3011;
Best Local Similarity 97.8%; Pred. No. 2.4e-86;
Matches 178; Conservative 3; Mismatches 1; Indels 0; Gaps 0;

QY 1 MAPITAYAAQOTRGLGCGIITSITGRKKNQVEGEVQIVSTAAQTFLATCINGVCMVTVYHGA 60
Db 1026 LAPITAYAAQOTRGLGCGIITSITGRKKNQVEGEVQIVSTAAQTFLATCINGVCMVTVYHGA 1085

QY 61 GRTTASPKGPVIOYNTYDKLVGWPAPQGSRLTPCTCGSSDLYLVTRHADVIPVRRR 120
Db 1086 GRTTASPKGPVIOYNTYDKLVGWPAPQGSRLTPCTCGSSDLYLVTRHADVIPVRRR 1145

QY 121 GDSRGLSPRPISYIKGSSGGPLLCAGHAGVIFRAAVCTRGVAKAVDFTIPVESLETTM 160
Db 1146 GDSRGLSPRPISYIKGSSGGPLLCAGHAGVIFRAAVCTRGVAKAVDFTIPVESLETTM 1255

QY 181 RS 182
Db 1206 RS 1207

```

```

RESULT 13
Q91RT4 PRELIMINARY: PRT; 181 AA.
AC Q91RT4;
DT 31-DEC-2001 (TrEMBLrel. 19, Created)
DT 31-DEC-2001 (TrEMBLrel. 19, Last sequence update)
DT 31-MAR-2003 (TrEMBLrel. 23, Last annotation update)
DE NS3 protease (Fragment).
OS Hepatitis C virus.
OC Viruses; ssRNA positive-strand viruses, no DNA stage: Flaviviridae;
OC Hepacivirus.
OX NCBI_TaxID=11103;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=PC.23;
RA Holland-Staley C.A., Kovari L.C., Golenberg E., Mayers D.L.;
RT "Genetic Diversity and response to IFN of the NS3 Protease Gene from
RT Clinical Strains of the Hepatitis C Virus.";
RL Submitted (APR-2001) to the EMBL/GenBank/DBJ databases.
DR EMBL: AF369219; AAK54544.1; -.
DR InterPro: IPR004109; HCV_NS3.
DR Pfam: PF02907; HCV_NS3; 1.
DR Protease.
KW NON_TER 1 1
FT NON_TER 181 181
SQ SEQUENCE 181 AA; 19059 MW; 1E53C47AE8B7E5C9 CRC64;

Query Match 98.4%; Score 938; DB 12; Length 181;
Best Local Similarity 98.3%; Pred. No. 9.6e-88;
Matches 178; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 2 APITAYAAQOTRGLGCGIITSITGRKKNQVEGEVQIVSTAAQTFLATCINGVCMVTVYHGA 61
Db 1 APITAYAAQOTRGLGCGIITSITGRKKNQVEGEVQIVSTAAQTFLATCINGVCMVTVYHGA 60

QY 62 TTTIASPKGPIVIOYNTYDKLVGWPAPQGSRLTPCTCGSSDLYLVTRHADVIPVRRR 121
Db 62 TTTIASPKGPIVIOYNTYDKLVGWPAPQGSRLTPCTCGSSDLYLVTRHADVIPVRRR 120

QY 122 DSRGSLSPRPISYIKGSSGGPLLCAGHAGVIFRAAVCTRGVAKAVDFTIPVESLETTM 181
Db 121 DSRGSLSPRPISYIKGSSGGPLLCAGHAGVIFRAAVCTRGVAKAVDFTIPVESLETTM 180

QY 182 S 182
Db 181 S 181

RESULT 14
Q91RS8 PRELIMINARY: PRT; 181 AA.
AC Q91RS8;
DT 01-DEC-2001 (TrEMBLrel. 19, Created)
DT 01-DEC-2001 (TrEMBLrel. 19, Last sequence update)
DT 01-MAR-2003 (TrEMBLrel. 23, Last annotation update)
DE NS3 protease (Fragment).
OS Hepatitis C virus.
OC Viruses; ssRNA positive-strand viruses, no DNA stage: Flaviviridae;
OC Hepacivirus.
OX NCBI_TaxID=11103;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=PC.176;
RA Holland-Staley C.A., Kovari L.C., Golenberg E., Mayers D.L.;
RT "Genetic Diversity and response to IFN of the NS3 Protease Gene from
RT Clinical Strains of the Hepatitis C Virus.";
RL Submitted (APR-2001) to the EMBL/GenBank/DBJ databases.
DR EMBL: AF369225; AAK54550.1; -.
DR InterPro: IPR004109; HCV_NS3.
DR Pfam: PF02907; HCV_NS3; 1.
DR Protease.
KW NON_TER 1 1
FT NON_TER 1 1

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Job time : 109 secs

FT NON_TER 181 181
SQ SEQUENCE 181 AA: 19114 MW: 574AC47AE8AEFFD2 CRC64:

Query Match 98.4%; Score 938; DB 12; Length 181;
Best Local Similarity 99.3%; Pred. No. 9.6e-88;
Matches 178; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 2 APITAYAQQRGLGCGIITSLTGROKNGVEGVIVSTANQFELATCINVGWCVTVHGAG 61
DB 1 APITAYAQQRGLGCGIITSLTGROKNGVEGVIVSTANQFELATCINVGWCVTVHGAG 60
CY 62 TRTIASPKGPVIQMYTNVCKDLVWKPAPQCSRLTCTCGSSDLYLVTRHADVTPVRRG 121
DB 61 TRTIASPKGPVIQMYTNVCKDLVWKPAPQARSLTCTCGSSDLYLVTRHADVTPVRRG 120
QY 122 DSRGSLSPRPISYLGSSGGPLLCFAGHAGVIFRAAVCTRGVAKAVDFIPVSLPTTMR 181
DB 121 DSRGSLSPRPISYLGSSGGPLLCFAGHAGVIFRAAVCTRGVAKAVDFIPVSLPTTMR 180
CY 182 S 182
DB 181 S 181

RESULT 15
Q91RT3 PRELIMINARY; PRI: 181 AA.
AC Q91RT3
DT 01-DEC-2001 (TrEMBLrel. 19, Created)
DI 01-DEC-2001 (TrEMBLrel. 19, Last sequence update)
DT 01-MAR-2003 (TrEMBLrel. 23, Last annotation update)
DE NS3 protease (Fragment).
OS Hepatitis C virus.
OC Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
OC Hepacivirus.
OX NCBI_TaxID=11103;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=Pt.11;
RA Holland-Staley C.A., Kovari L.C., Golenborg E., Kayors D.L.;
RT "Genetic Diversity and Response to IFN of the NS3 Protease Gene from
Clinical Strains of the Hepatitis C Virus."
RL Submitted (AFR-2001) to the EMBL/GenBank/DBJ databases.
DR EMBL: AF369220; AAK54545.1;
DR InterPro: IPR004109; HCV_NS3;
DR Pfam: PF02907; HCV_NS3; 1.
KW Protease.
KW NON_TER 181 181
FT NON_TER 181 181
SQ SEQUENCE 181 AA: 19116 MW: 6648807F49B1D33 CRC64:

Query Match 98.4%; Score 938; DB 12; Length 181;
Best Local Similarity 98.3%; Pred. No. 4.6e-85;
Matches 178; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 2 APITAYAQQRGLGCGIITSLTGROKNGVEGVIVSTANQFELATCINVGWCVTVHGAG 61
DB 1 APITAYAQQRGLGCGIITSLTGROKNGVEGVIVSTANQFELATCINVGWCVTVHGAG 60
CY 62 TRTIASPKGPVIQMYTNVCKDLVWKPAPQCSRLTCTCGSSDLYLVTRHADVTPVRRG 121
DB 61 TRTIASPKGPVIQMYTNVCKDLVWKPAPQARSLTCTCGSSDLYLVTRHADVTPVRRG 120
QY 122 DSRGSLSPRPISYLGSSGGPLLCFAGHAGVIFRAAVCTRGVAKAVDFIPVSLPTTMR 181
DB 121 DSRGSLSPRPISYLGSSGGPLLCFAGHAGVIFRAAVCTRGVAKAVDFIPVSLPTTMR 180
CY 182 S 182
DB 181 S 181

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: September 26, 2003, 22:20:58 : Search time 72 seconds

(without alignments)
370,362 Million cell updates/sec

Title: US-09-965-594-1

Perfect score: 953

Sequence: 1 NAFITAYAGTGTGACGIICT.....GVAKAYGFIIVRSLEITWMS 182

Scoring Table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 1:07863 seqs, 158726573 residues

Total number of hits satisfying chosen parameters: 110762

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 94

Maximum Match 94

Listing first 45 summaries

Database : A_Geneseq_19Jun03.*

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2: /SIDSL/gcgdata/geneseq/geneseq-emb1/AA1981.DAT.*
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5: /SIDSL/gcgdata/geneseq/geneseq-emb1/AA1984.DAT.*
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10: /SIDSL/gcgdata/geneseq/geneseq-emb1/AA1989.DAT.*
11: /SIDSL/gcgdata/geneseq/geneseq-emb1/AA1990.DAT.*
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13: /SIDSL/gcgdata/geneseq/geneseq-emb1/AA1992.DAT.*
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19: /SIDSL/gcgdata/geneseq/geneseq-emb1/AA1998.DAT.*
20: /SIDSL/gcgdata/geneseq/geneseq-emb1/AA1999.DAT.*
21: /SIDSL/gcgdata/geneseq/geneseq-emb1/AA2000.DAT.*
22: /SIDSL/gcgdata/geneseq/geneseq-emb1/AA2001.DAT.*
23: /SIDSL/gcgdata/geneseq/geneseq-emb1/AA2002.DAT.*
24: /SIDSL/gcgdata/geneseq/geneseq-emb1/AA2003.DAT.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	946	99.3	586	AAE18689	HCV-1 NS3/4a mutant
2	946	99.3	586	AAE18689	Hepatitis C virus
3	946	99.3	586	AAE18689	HCV-1 NS3/4a confo
4	944	99.1	3011	AAE18689	HCV genomic amino
5	943	99.0	609	AAE18689	Hepatitis C virus
6	943	99.0	1766	AAE18689	Sequence encoded i
7	943	99.0	1766	AAE18689	Protein: sequence o
8	943	99.0	2261	AAE18689	Peptide encoded by
9	943	99.0	2301	AAE18689	Sequence encoded i

10	943	99.0	2436	10	AAE18689	Sequence encoded i
11	943	99.0	2436	10	AAE18689	Peptide encoded by
12	943	99.0	2772	21	AAE18689	Protein: encoded by
13	943	99.0	2816	14	AAE18689	HCV-1 polyprotein
14	943	99.0	2894	16	AAE18689	Composite hepatitis
15	943	99.0	2955	20	AAE18689	Amino acid sequenc
16	943	99.0	2955	21	AAE18689	Polyprotein encode
17	943	99.0	3011	13	AAE18689	Compiled HCV sequ
18	943	99.0	3011	14	AAE18689	Hepatitis C virus
19	943	99.0	3011	17	AAE18689	Hepatitis C virus
20	943	99.0	3011	18	AAE18689	HCV polyprotein
21	943	99.0	3011	19	AAE18689	HCV polyprotein
22	943	99.0	3011	23	AAE18689	Hepatitis C virus
23	943	99.0	3011	23	AAE18689	HCV polyprotein la
24	942	98.8	632	23	AAE18689	Hepatitis C virus
25	942	98.8	632	23	AAE18689	Hepatitis C virus
26	942	98.8	686	23	AAE18689	Hepatitis C virus
27	942	98.8	686	23	AAE18689	Hepatitis C virus
28	942	98.8	686	23	AAE18689	Hepatitis C virus
29	942	98.8	686	23	AAE18689	Hepatitis C virus
30	942	98.8	686	23	AAE18689	Hepatitis C virus
31	942	98.8	686	23	AAE18689	Hepatitis C virus
32	942	98.8	686	23	AAE18689	Hepatitis C virus
33	942	98.8	686	23	AAE18689	Hepatitis C virus
34	942	98.8	686	23	AAE18689	Hepatitis C virus
35	942	98.8	686	23	AAE18689	Hepatitis C virus
36	942	98.8	686	23	AAE18689	Hepatitis C virus
37	942	98.8	686	23	AAE18689	Hepatitis C virus
38	942	98.8	686	23	AAE18689	Hepatitis C virus
39	942	98.8	686	23	AAE18689	Hepatitis C virus
40	942	98.8	686	23	AAE18689	Hepatitis C virus
41	942	98.8	686	23	AAE18689	Hepatitis C virus
42	942	98.8	686	23	AAE18689	Hepatitis C virus
43	942	98.8	686	23	AAE18689	Hepatitis C virus
44	942	98.8	686	23	AAE18689	Hepatitis C virus
45	942	98.8	686	23	AAE18689	Hepatitis C virus

ALIGNMENTS

RESULT 1

AAE18689
10 AAE18689 standard: Protein; 686 AA.
XX
XX AAE18689:
AC
XX
XX 17-MAY-2002 (first entry)
XX
XX HCV-1 NS3/4a mutant conformational antigen.
XX
XX Hepatitis C virus: NS3/4a antigen; HCV infection; mutant; mutin.
XX
XX Hepatitis C virus type 1.
XX
XX Synthetic.

Key Location/Qualifiers
FI Misc-difference 403 /note- "Wild type Thr substituted with Pro"
FI Misc-difference 404 /note- "Wild type Ser substituted with Ile"
XX
XX WC200196675-A2.
XX
XX 20-DEC-2001.
XX
XX 14-JUN-2001; 2001WO-0519369.
XX
XX 15-JUN-2000; 2000US-212082P.
XX
XX 02-APR-2001; 2001US-280811P.
XX
XX 02-APR-2001; 2001US-280867P.
XX
XX (CHIR) CHIRON CORP.

```

XX Chien DY, Arcangel P, Tandeske L, George-Nascimento C, Coit D;
PI Medina-Selby A;
XX WP1: 2002-179522/23.
XX DR N-PSDB; AAD29795.
XX PT
XX PT Immunassay solid support useful for detecting hepatitis C virus
XX PT infection in a biological sample, comprises at least one of HCV
XX PT anti-core antibody and HCV NS3/4a epitope, bound to the support.
XX PS Example 2; Fig 4; 87pp; English.
XX CC The present invention relates to hepatitis C virus (HCV) core antigen
XX CC and NS (nonstructural) 3/4a antibody combination assay that can detect
XX CC both HCV antigens and antibodies present in a sample using a single
XX CC solid matrix as well as immunoassay solid supports for use in the assay.
XX CC The solid support is useful for detecting HCV infection in a biological
XX CC sample. The present sequence is HCV-1 NS3/4a mutant conformational
XX CC antigen. This sequence is used in the exemplification of the invention.
XX SQ Sequence 686 AA;
XX Query Match 99.3%; Score 946; DA 23; Length 686;
XX Best Local Similarity 98.9%; Pred. No. 1.2e-89;
XX Matches 180; Conservative 2; Mismatches 0; Indels 0; Gaps 0;
QY 1 MAPITAYAQTRGLGCIITSLTGKDNQVEGEVQVSTAAQTFLATCINGVCTVYHGA 60
DB 1 MAPITAYAQTRGLGCIITSLTGKDNQVEGEVQVSTAAQTFLATCINGVCTVYHGA 60
QY 61 GTRIASPKGPVIOMYINVDKLVGNWPAPOGSRSLTPTCGSSDLYLVTTHADVIPVRRR 120
DB 61 GTRIASPKGPVIOMYINVDKLVGNWPAPOGSRSLTPTCGSSDLYLVTTHADVIPVRRR 120
QY 121 GDSRGSLLSPRPISYLGKSSGGPILCPAGHAVGIFRAAVCTRGVAKAVDFIPVESLETTM 180
DB 121 GDSRGSLLSPRPISYLGKSSGGPILCPAGHAVGIFRAAVCTRGVAKAVDFIPVENLETTM 180
QY 181 RS 182
DB 181 RS 182
XX RESULT 3
XX ABG72261
XX ID ARG72261 standard; Protein: 686 AA.
XX AC ABG72261;
XX DT 06-MAR-2003 (first entry)
XX DE HCV-1 NS3/4a conformational antigen.
XX

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XX 15-JUN-2000; 2000US-212082P.
XX 02-APR-2001; 2001US-280811P.
XX 02-APR-2001; 2001US-280867P.
XX (CHIR) CHIRON CORP.
XX Chien DY, Arcangel P, Tandeske L, George-Nascimento C, Coit D;
PI Medina-Selby A;
XX WP1: 2002-090228/12.
XX DR N-PSDB; ABK15344.
XX PT
XX PT Immunassay solid support, useful for detecting hepatitis C virus
XX PT infection in biological sample, comprises HCV NS3/4a conformational
XX PT epitope and multiple epitope fusion antigen bound to the support.
XX PS Claim 5; Fig 3; 92pp; English.
XX CC The present invention relates to a new immunoassay solid support
XX CC consisting essentially of at least one hepatitis C virus (HCV) NS3/4a
XX CC conformational epitope and a multiple epitope fusion antigen (MEFA),
XX CC bound to the support. The NS3/4a conformational epitope and/or
XX CC MEFA reacts specifically with anti-HCV antibodies present in a biological
XX CC sample from an HCV-infected individual. The immunoassay of the invention
XX CC is useful for detecting hepatitis C virus infection in a biological
XX CC sample. The method of the invention provides a sensitive, accurate
XX CC diagnostic and prognostic tool to provide adequate patient care and to
XX CC prevent transmission of HCV by blood and by blood products, or by
XX CC personal contact. Use of NS3/4a conformational epitope in combination
XX CC with MEFA, provides a sensitive and reliable method for detecting early
XX CC HCV seroconversion. Use of MEFA has the added advantages of decreasing
XX CC masking problems, improving sensitivity in detecting antibodies by
XX CC allowing a greater number of epitopes on a unit surface area of
XX CC substrate, and improving substrate. Detection accuracy is increased and
XX CC the incidence of false results is reduced because of the identification
XX CC and the use of highly immunogenic HCV antigens which are present during
XX CC the early stages of HCV seroconversion. The present amino acid sequence
XX CC represents the non-structural protein NS3/4a conformational epitope of
XX CC the invention.
XX SQ Sequence 686 AA;
XX Query Match 99.3%; Score 946; DA 23; Length 686;
XX Best Local Similarity 98.9%; Pred. No. 1.2e-89;
XX Matches 180; Conservative 2; Mismatches 0; Indels 0; Gaps 0;
QY 1 MAPITAYAQTRGLGCIITSLTGKDNQVEGEVQVSTAAQTFLATCINGVCTVYHGA 60
DB 1 MAPITAYAQTRGLGCIITSLTGKDNQVEGEVQVSTAAQTFLATCINGVCTVYHGA 60
QY 61 GTRIASPKGPVIOMYINVDKLVGNWPAPOGSRSLTPTCGSSDLYLVTTHADVIPVRRR 120
DB 61 GTRIASPKGPVIOMYINVDKLVGNWPAPOGSRSLTPTCGSSDLYLVTTHADVIPVRRR 120
QY 121 GDSRGSLLSPRPISYLGKSSGGPILCPAGHAVGIFRAAVCTRGVAKAVDFIPVESLETTM 180
DB 121 GDSRGSLLSPRPISYLGKSSGGPILCPAGHAVGIFRAAVCTRGVAKAVDFIPVENLETTM 180
QY 181 RS 182
DB 181 RS 182
XX RESULT 3
XX ABG72261
XX ID ARG72261 standard; Protein: 686 AA.
XX AC ABG72261;
XX DT 06-MAR-2003 (first entry)
XX DE HCV-1 NS3/4a conformational antigen.
XX

```

Immunosassay solid support; Hepatitis C Virus type-1; HCV-1;
NS3/4a conformational epitope; multiple epitope fusion antigen;
MEFA; anti-HCV antibody; NS3/4a conformational antigen;
HCV infection; mutant; muten.

Hepatitis C virus type 1.
Synthetic.

Key Location/Qualifiers

Region 2...586
/note= "Corresponds to amino acid residues 1027-1171
of HCV-1 NS3/4a polypeptide"

Misc-difference 403

/note= "Substitution of wild-type Thr to Pro"

Misc-difference 404

/note= "Substitution of wild-type Ser to Leu"

US2002146685-A1.

10-OCT-2002.

14-JUN-2001; 2001US-0861654.

15-JUN-2000; 2000US-2120827.

02-APR-2001; 2001US-2808112.

02-APR-2001; 2001US-2808672.

(CHIEF) CHIEN D Y.

(ARCA) ARCANDEL P.

(TAND) TANDESKE L.

(GEOR) GEORGE-NASCIMENTO C.

(COLT) COLT D.

(MEDT) MEDINA-SELBY A.

Chien DY, Arcangel P, Tandeske L, George-Nascimento C, Colt D;
Medina-Selby A;

WPI: 2003-147573/14.

N-PSDB; ARX14410.

Immunosassay solid support for detecting Hepatitis C Virus infection in
biological samples, comprises Hepatitis C Virus conformational epitope
and multiple epitope fusion antigen.

Claim 2: Fig 3A-3D; 45pp; English.

The present invention relates to immunoassays comprising Hepatitis C
virus (HCV) NS3/4a conformational epitope and multiple epitope fusion
antigen (MEFA), bound to a solid support. The NS3/4a epitope and/or
the multiple epitope fusion antigen react with anti-HCV antibodies
present in a biological sample from an HCV-infected individual. The
immunoassays and methods of the invention are useful for detecting
HCV infection in a biological sample. The inventive immunoassay solid
support provides a sensitive and reliable method for detecting early
HCV seroconversion. The assays can detect HCV infection caused by any
six known genotypes of HCV. The use of the multiple epitope fusion
proteins decreases masking problems, improves sensitivity in detecting
antibodies by allowing a greater number of epitopes on a unit area
of substrate, and improves selectivity. The present sequence
represents HCV type 1 (HCV-1) NS3/4a conformational antigen, a mutant
of the HCV-1 NS3/4a polypeptide.

Sequence 686 AA;

Query Match 99.3%; Score 346; DR 24; Length 686;

Best Local Similarity 98.9%; Pred. No. 1.2e-86;

Matches 180; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

1 MAPITAYAOQTGRLGCIITSLTGRKNOVEGEVQIVSTAAQTFLATCINGVCTVYHGA 60
|||||
1 MAPITAYAOQTGRLGCIITSLTGRKNOVEGEVQIVSTAAQTFLATCINGVCTVYHGA 60

61 GTTITASPKGPVQMTYTNVDKLVGWPAPOGGRSITPTCGSSDLYLVTRHADVIPVRR 120

DB 61 GTTITASPKGPVQMTYTNVDKLVGWPAPOGGRSITPTCGSSDLYLVTRHADVIPVRR 120
QY 121 GDSKGSLLSPRP:SYLKSGSGGPGLLCPAGHAGVCFRAAVCTIRGVAKAVDFIPVESLETTM 180
|||||
DB 121 GDSKGSLLSPRP:SYLKSGSGGPGLLCPAGHAGVCFRAAVCTIRGVAKAVDFIPVENLETTM 180

QY 181 RS 182

DB 181 RS 182

RES:LT_4

AA44C120

AA44C120 standard; Protein: 3011 AA.

AC AAR40120;

25-MAR-2003 (updated)

27-JAN-1994 (first entry)

HCV genomic amino acid sequence isolated from infected human LG.

Hepatitis C Virus; Non-A, non-B Hepatitis Virus; HCV; NANBHV;

human growth hormone; HGH; secretion signal; fusion protein;

vacccine.

Hepatitis C Virus.

MO9315193-A1.

05-AUG-1993.

29-JAN-1993; 93WO-0500907.

31-JAN-1992; 92US-0630C24.

(ABBO) ABBOTT LAB.

Bode SL, Casey JM, Desai SM, Devare SG, Frail DE;
Yamaguchi J, Zeck BJ;

WPI: 1993-258673/32.

New plasmid pHCV-162 is a mammalian expression systems for HCV
proteins - useful for diagnosing HCV infection and as vaccines
for preventing HCV infection
Example 1; Page 39-49; 100pp; English.

RNA was isolated from the plasma of a HCV seropositive human
(designated "IG") and cDNA was prepared from it. The cDNA was
PCR amplified using specific primers with sequences based
on the prototype HCV-1 cDNA sequence (GENBANK M62321). Further
amplification using nested primers resulted in 7 adjacent HCV DNA
fragments which could be assembled into a full-length sequence. The
DNA sequence was determined and translated into the genomic amino
acid sequence. Comparison of the LG genomic amino acid sequence
with that from HCV-1 showed 134 amino acid differences.
(Updated on 25-MAR-2003 to correct PN field.)

Sequence 3011 AA;

Query Match 99.1%; Score 944; DR 14; Length 3011;

Best Local Similarity 98.4%; Pred. No. 1.4e-88;

Matches 179; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

1 MAPITAYAOQTGRLGCIITSLTGRKNOVEGEVQIVSTAAQTFLATCINGVCTVYHGA 60

1026 LAPITAYAOQTGRLGCIITSLTGRKNOVEGEVQIVSTAAQTFLATCINGVCTVYHGA 1085

61 GTTITASPKGPVQMTYTNVDKLVGWPAPOGGRSITPTCGSSDLYLVTRHADVIPVRR 120
|||||

DB	1086	GIHTIASPKPVQMTNVPDSLVGVNPAFGARSLPCTCGSSDLVLTTHADVIVRRR	1144
OY	121	GDGRGSLSPRPISYLYKSGGGHLLCPAGHAGVIFRAAVCTRGVAKAVDFIPVESLETTM	180
DB	1146	GDGRGSLSPRPISYLYKSGGGHLLCPAGHAGVIFRAAVCTRGVAKAVDFIPVESLETTM	1205
OY	181	RS 182	
DB	1206	RS 1207	
RESULT 5			
AAR51170			
ID	AAR51170	standard; peptide: 609 AA.	
XX	AAR51170;		
XX	20-OCT-1994	(first entry)	
XX	Hepatitis C virus non-structural protein 3.		
DE			
XX	Peptide; antibody: hepatitis C virus; HCV; identification;		
KW	diagnosis; non-A non-B hepatitis; NANB; detection.		
XX			
OS	Hepatitis C virus.		
XX			
PN	JP06056891-A.		
XX			
PD	01-MAR-1994.		
XX			
PF	05-AUG-1992; 92JP-0209201.		
XX			
PR	05-AUG-1992; 92JP-0209201.		
XX			
PA	(OLYU) OLYMPUS OPTICAL CO LTD.		
XX			
DR	WPI: 1994-106863/13.		
XX			
PT	New peptide(s) reactive with anti-hepatitis C virus antibody .		
PT	For specific, early diagnosis of HCV infection		
XX			
PS	Disclosure; Page 9-10; 15pp; Japanese.		
XX			
CC	Peptide fragments of the non-structural protein (NS) are reactive		
CC	with and can detect antibodies against the NS3 domain of HCV. The		
CC	peptides can be used for diagnosis of HCV infection. Non-specific		
CC	reaction can be inhibited and misdiagnosis of HCV infection can be		
CC	decreased. See AAR51162-70.		
XX			
SO	Sequence 609 AA;		
Query Match	59.0%; Score 943; DB 15; Length 609;		
Best Local Similarity	98.4%; Pred. No. 2,16-80;		
Matches	179; Conservative 3; Mismatches 0; Indels 0; Gaps 0;		
OY	1	MAHITAYAQCTRGELLCILSLTGROKNQVESHQVLSAACTFLATCINGVCTVYHGA	60
DB	20	LAPITAYAQCTRGELLCILSLTGROKNQVESVQVLSAACTFLATCINGVCTVYHGA	79
OY	61	GIHTIASPKPVQMTNVPDSLVGVNPAFGARSLPCTCGSSDLVLTTHADVIVRRR	120
DB	80	GIHTIASPKPVQMTNVPDSLVGVNPAFGARSLPCTCGSSDLVLTTHADVIVRRR	139
OY	121	GDGRGSLSPRPISYLYKSGGGHLLCPAGHAGVIFRAAVCTRGVAKAVDFIPVESLETTM	180
DB	140	GDGRGSLSPRPISYLYKSGGGHLLCPAGHAGVIFRAAVCTRGVAKAVDFIPVENLETTM	199
OY	181	RS 182	
DB	200	RS 201	

RESULT 7
AAP90158
ID AAP90158 standard; protein: 1786 AA.
XX AC AAP90158:
XX
XX 25-MAR-2003 (updated)
DT 10-NOV-1989 (first entry)
XX
XX Protein sequence of hepatitis C virus composite cDNA.
XX
XX Hepatitis C virus; vaccine.
XX
XX Pan troglodytes.
XX
XX GB2212511-A.
XX
XX 26-JUL-1989.
XX
XX 18-NOV-1988; 88GB-0027024.
XX
XX 18-NOV-1987; 87US-0122714.
PR 30-DEC-1987; 87US-0139886.
PR 26-FEB-1988; 88US-0161072.
PR 26-OCT-1988; 88US-0263584.
XX
XX (CHIR) CHIRON CORPORATION.
XX
XX Houghton M, Choo QL, Kuo G;
XX
XX W21; 1989-215054/30.
DR N-PSDB; AAN90327.
XX
XX Hepatitis C virus gene - used for prodn. of polynucleotide probes.
PT polypeptide(s) and antibodies for diagnosis, prevention and treatment
PT of infection.
XX
XX Disclosure; fig 26; 30pp; English.

PS The sequence is encoded by the composite cDNA of AAN90327. These
CC antigens react with antibodies in patients with non-A non-B hepatitis
CC (NANBH). They can be used to diagnose HCV-induced NANBH, to raise
CC antibodies for immunoassay or treatment, or to produce vaccines.
CC (Updated on 25-MAR-2003 to correct PR field.)
XX
XX Sequence 1786 AA;
SQ

Query Match 99.0%; Score 943; DB 10; Length 1786;
Best Local Similarity 98.4%; Pred. No. 8.9e-89;
Matches 179; Conservative 3; Mismatches 0; Indels 0; Gaps 0;
QY 1 MAPITAYAOQTRGLGCGIITSLTGRDNKQVGEVQIVSTAAQTFLATCINGVQWTVYHGA 60
DB 310 LAPITAYAOQTRGLGCGIITSLTGRDNKQVGEVQIVSTAAQTFLATCINGVQWTVYHGA 369
QY 61 GTRTIASPKGPVQMYTNVDKDLVGNWPAQGSRLTPTCGSSDLYLVTRHADVIPVRR 120
DB 370 GTRTIASPKGPVQMYTNVDKDLVGNWPAQGSRLTPTCGSSDLYLVTRHADVIPVRR 429
QY 121 GDSRGSLLSPRPISYLGKSGSGPLCPAGHAGVIFRAAVCTRGVAKAVDFIPVESLETTM 180
DB 430 GDSRGSLLSPRPISYLGKSGSGPLCPAGHAGVIFRAAVCTRGVAKAVDFIPVENLETTM 489
QY 181 RS 182
DB 490 RS 491

RESULT 8
AAP90164
ID AAP90164 standard; protein: 2261 AA.
XX
XX AAP90164:

XX 25-MAR-2003 (updated)
DT 10-NOV-1989 (first entry)
XX
XX Peptide encoded by composite hepatitis C virus cDNA.
XX
XX Hepatitis C virus; clone 121; clone 150; probe; vaccine.
XX
XX Pan troglodytes.
XX
XX GB2212511-A.
XX
XX 26-JUL-1989.
XX
XX 18-NOV-1988; 88GB-0027024.
XX
XX 18-NOV-1987; 87US-0122714.
PR 30-DEC-1987; 87US-0139886.
PR 26-FEB-1988; 88US-0161072.
PR 26-OCT-1988; 88US-0263584.
XX
XX (CHIR) CHIRON CORPORATION.
XX
XX Houghton M, Choo QL, Kuo G;
XX
XX W21; 1989-215054/30.
DR N-PSDB; AAN90331.
XX
XX Hepatitis C virus gene - used for prodn. of polynucleotide probes,
PT polypeptide(s) and antibodies for diagnosis, prevention and
PT treatment of infection.
XX
XX Disclosure; fig 32; 235pp; English.
XX
XX The sequence is the peptide encoded by the composite hepatitis C
CC virus (HCV) cDNA of AAN90331. The polypeptides are used to diagnose
CC HCV-induced NANBH, to raise antibodies for immunoassay or treatment,
CC or to produce vaccines.
CC (Updated on 25-MAR-2003 to correct PR field.)
XX
XX Sequence 2261 AA;
SQ

Query Match 99.0%; Score 943; DB 10; Length 2261;
Best Local Similarity 98.4%; Pred. No. 1.2e-88;
Matches 179; Conservative 3; Mismatches 0; Indels 0; Gaps 0;
QY 1 MAPITAYAOQTRGLGCGIITSLTGRDNKQVGEVQIVSTAAQTFLATCINGVQWTVYHGA 60
DB 401 LAPITAYAOQTRGLGCGIITSLTGRDNKQVGEVQIVSTAAQTFLATCINGVQWTVYHGA 460
QY 61 GTRTIASPKGPVQMYTNVDKDLVGNWPAQGSRLTPTCGSSDLYLVTRHADVIPVRR 120
DB 461 GTRTIASPKGPVQMYTNVDKDLVGNWPAQGSRLTPTCGSSDLYLVTRHADVIPVRR 520
QY 121 GDSRGSLLSPRPISYLGKSGSGPLCPAGHAGVIFRAAVCTRGVAKAVDFIPVESLETTM 180
DB 521 GDSRGSLLSPRPISYLGKSGSGPLCPAGHAGVIFRAAVCTRGVAKAVDFIPVENLETTM 580
QY 181 RS 182
DB 581 RS 582

RESULT 9
AAP92047
ID AAP92047 standard; protein: 2301 AA.
XX
XX AAP92047:

XX 25-MAR-2003 (updated)
DT 02-MAR-1990 (first entry)
XX
XX Sequence encoded in the hepatitis C virus (HCV) cDNA inserts in clones

```

DE 12f through 15e.
XX
XX Hepatitis C virus (HCV): non-A, non-B hepatitis (HANBH).
KW
XX
XX Hepatitis C virus.
OS
XX
XX Hepatitis C virus.
PN
XX
XX EP318216-A.
XX
XX 31-MAY-1989.
XX
XX
XX 18-NOV-1988; 88EP-0310922.
XX
XX
XX 18-NOV-1987; 87US-C122714.
XX
XX 30-DEC-1987; 87US-C139886.
XX
XX 26-FEB-1988; 88US-0161072.
XX
XX 06-MAY-1988; 88US-0191263.
XX
XX 26-OCT-1988; 88US-0263584.
XX
XX 14-NOV-1988; 88US-0272450.
XX
XX (CHIR ) CHIRON CORP.
XX
XX Houghton M, Choo QL, Kuo G;
XX
XX WPI; 1989-159274/22.
XX
XX N-PSDB; AAN92103.
XX
XX Purified hepatitis C virus
XX
XX - and associated nucleic acids and polypeptide(s)
XX
XX Claim 13; Figure 32-1 - 32-7; 139 pp; English.
XX
XX It is the sequence encoded in the open reading frame of hepatitis C virus
XX
XX (HCV) cDNA inserts in clones 12f through 15e. It is antigenic and could
XX
XX be used in immunoassay reagents and vaccines and to generate antibodies
XX
XX useful in diagnosis and passive immunotherapy for HCV infection/non-A,
XX
XX non-B hepatitis.
XX
XX (Updated on 25-MAR-2003 to correct PR field.)
XX
XX (Updated on 25-MAR-2003 to correct PI field.)
XX
XX
XX
XX Sequence 2301 AA;
XX
XX Query Match 99.0%; Score 943; DB 10; Length 2301;
XX
XX Best Local Similarity 98.4%; Pred. No. 1.2e-88;
XX
XX Matches 179; Conservative 3; Mismatches 0; Indels 0; Gaps 0;
XX
XX QY 1 MAPITAYAOOTRGLGCIITSLTGRDKNQVEGEVCIYSTAAQTFLACINGVCTVYHGA 60
XX
XX DE 401 LAPITAYAOOTRGLGCIITSLTGRDKNQVEGEVCIYSTAAQTFLACINGVCTVYHGA 460
XX
XX QY 61 GTRTIASPKGPVIOQMTNVKDLVGNWPAQGSRLTPTCGSSDLYLVTRHADVIPVRRR 120
XX
XX DB 461 GTRTIASPKGPVIOQMTNVKDLVGNWPAQGSRLTPTCGSSDLYLVTRHADVIPVRRR 520
XX
XX QY 121 GDSKGSLLSPRPISYLVKGSGLGPGAGHAGVIFRAAVCTRGVAKAVDFIPVESLETTM 180
XX
XX DB 521 GDSKGSLLSPRPISYLVKGSGLGPGAGHAGVIFRAAVCTRGVAKAVDFIPVENLETTM 580
XX
XX QY 181 RS 182
XX
XX DB 561 RS 582
XX
XX
XX RESULT 10
XX
XX AAP92050
XX
XX ID AAP92050 standard; protein: 2436 AA.
XX
XX
XX AAP92050;
XX
XX AC
XX
XX AC
XX
XX 25-MAR-2003 (updated)
XX
XX DT 02-MAR-1990 (first entry)
XX
XX
XX Sequence encoded in the hepatitis C virus (HCV) cDNA inserts in clones
XX
XX K9-1 through 15e.

```

```

XX
XX Hepatitis C virus (HCV): non-A, non-B hepatitis (HANBH)
XX
XX Hepatitis C virus.
XX
XX PN
XX
XX EP318216-A.
XX
XX 31-MAY-1989.
XX
XX
XX 18-NOV-1988; 88EP-0310922.
XX
XX
XX 18-NOV-1987; 87US-C122714.
XX
XX 30-DEC-1987; 87US-C139886.
XX
XX 26-FEB-1988; 88US-0161072.
XX
XX 06-MAY-1988; 88US-0191263.
XX
XX 26-OCT-1988; 88US-0263584.
XX
XX 14-NOV-1988; 88US-0271450.
XX
XX (CHIR ) CHIRON CORP.
XX
XX Houghton M, Choo QL, Kuo G;
XX
XX WPI; 1989-159274/22.
XX
XX N-PSDB; AAN92106.
XX
XX Purified hepatitis C virus
XX
XX - and associated nucleic acids and polypeptide(s)
XX
XX Claim 13; Figure 47-1 - 47-8; 139 pp; English.
XX
XX It is the sequence encoded in the open reading frame of hepatitis C virus
XX
XX (HCV) cDNA inserts in clones K9-1 through 15e. It is antigenic and could
XX
XX be used in immunoassay reagents and vaccines and to generate antibodies
XX
XX useful in diagnosis and passive immunotherapy for HCV infection/non-A,
XX
XX non-B hepatitis.
XX
XX (Updated on 25-MAR-2003 to correct PR field.)
XX
XX (Updated on 25-MAR-2003 to correct PI field.)
XX
XX
XX
XX Sequence 2436 AA;
XX
XX Query Match 99.0%; Score 943; DB 10; Length 2436;
XX
XX Best Local Similarity 98.4%; Pred. No. 1.3e-86;
XX
XX Matches 179; Conservative 3; Mismatches 0; Indels 0; Gaps 0;
XX
XX QY 1 MAPITAYAOOTRGLGCIITSLTGRDKNQVEGEVCIYSTAAQTFLACINGVCTVYHGA 60
XX
XX DB 576 LAPITAYAOOTRGLGCIITSLTGRDKNQVEGEVCIYSTAAQTFLACINGVCTVYHGA 635
XX
XX QY 61 GTRTIASPKGPVIOQMTNVKDLVGNWPAQGSRLTPTCGSSDLYLVTRHADVIPVRRR 120
XX
XX DB 636 GTRTIASPKGPVIOQMTNVKDLVGNWPAQGSRLTPTCGSSDLYLVTRHADVIPVRRR 695
XX
XX QY 121 GDSKGSLLSPRPISYLVKGSGLGPGAGHAGVIFRAAVCTRGVAKAVDFIPVESLETTM 180
XX
XX DB 696 GDSKGSLLSPRPISYLVKGSGLGPGAGHAGVIFRAAVCTRGVAKAVDFIPVENLETTM 755
XX
XX QY 181 RS 182
XX
XX DB 756 RS 757
XX
XX
XX RESULT 11
XX
XX AAP90288
XX
XX ID AAP90288 standard; protein: 2436 AA.
XX
XX
XX AAP90288;
XX
XX AC
XX
XX AC
XX
XX 25-MAR-2003 (updated)
XX
XX DT 19-JUL-2001 (updated)
XX
XX DT 01-NOV-1989 (first entry)
XX
XX
XX Peptide encoded by composite hepatitis C cDNA.
XX

```


KW Hepatitis C virus; clone 15c; clone kg-1; probe: vaccine.

XX Pan troglodytes.

XX GB2212911-A.

XX 26-JUL-1989.

XX 18-NOV-1988; 583B-0027224.

XX 19-NOV-1987; 87US-0122714.

XX 30-DEC-1987; 87US-0139886.

XX 26-FEB-1986; 88US-0161072.

XX 25-OCT-1988; 88US-0263584.

XX (CHIR) CHIRON CORPORATION.

XX Houghton M, Choo QL, Kuo G;

XX WPI; 1989-215054/30.

XX N-PSDB; AAN9336.

XX Hepatitis C virus gene - used for prodn. of polynucleotide protein.

XX polypeptide(s) and antibodies for diagnosis, prevention and

XX treatment of infection.

XX Disclosure; fig 47-1 to 47-8; 235pp; English.

XX The sequence is the peptide encoded by the composite hepatitis C

XX virus (HCV) cDNA of AAN9336. The polypeptides are used to

XX diagnose HCV-induced NANBH. To raise antibodies for

XX immunoassay or treatment, or to produce vaccines.

XX (N.B. This record was resubmitted to correct errors in the sequence.)

XX (Updated on 25-MAR-2003 to correct PR field.)

XX Sequence 2436 AA;

XX Query Match 99.0%; Score 943; DB 10; Length 2436;

XX Best Local Similarity 98.4%; Prod. No. 1.3e-88;

XX Matches 179; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 MAPITAYAOQTGRLGCGIITSLTGRDNKQVEGEVQIVSTAAQTFLATCINGVCTVYHGA 60

Db 576 LAPITAYAOQTGRLGCGIITSLTGRDNKQVEGEVQIVSTAAQTFLATCINGVCTVYHGA 635

QY 61 GTTITASPQGPVQIOMYTNVDKLVGWPAPQGSRLTPTCTCGSSDLVYVTRHADVIPVRRR 120

Db 636 GTTITASPQGPVQIOMYTNVDKLVGWPAPQGSRLTPTCTCGSSDLVYVTRHADVIPVRRR 695

QY 121 GDSRGLLSPPRISYLGSGGSPILCPAGHAGVIFRAAVCTRGVAKAVDFIPVESLETTM 180

Db 696 GDSRGLLSPPRISYLGSGGSPILCPAGHAGVIFRAAVCTRGVAKAVDFIPVESLETTM 755

QY 181 RS 182

Db 756 RS 757

RESULT 12

AAB18540

ID AAB18540 standard; Protein: 2772 AA.

XX AAB18540;

XX 15-JAN-2001 (first entry)

XX Protein encoded by a cDNA compiled Hepatitis C virus cDNA clones.

XX Hepatitis C virus; HCV; antisense polynucleotide; polypeptide;

XX viral infectivity; viral replication.

XX Hepatitis C virus.

XX

PN EF1034785-A2.

XX 13-SEP-2000.

XX 16-MAR-1990; 2000EP-0109602.

XX 17-MAR-1989; 89US-0325338.

XX 20-APR-1989; 89US-0341334.

XX 18-MAY-1989; 89US-0355002.

XX 16-MAR-1990; 90EP-0302866.

XX (CHIR) CHIRON CORP.

XX Houghton M, Choo Q, Kuo G;

XX WPI; 2000-566891/53.

XX N-PSDB; AAN75296.

XX Novel composition comprising a hepatitis C virus antisense

XX polynucleotide which is complementary to or corresponds to a sense

XX strand of the virus genome, and selectively hybridizes to it.

XX Example; Fig 16; 75pp; English.

XX The specification describes a pharmaceutical composition which

XX comprises a hepatitis C virus (HCV) antisense polynucleotide. The

XX HCV is characterized by a positive stranded RNA genome which has

XX 40% homology at the polypeptide level to a HCV polyprotein. The

XX antisense polynucleotide binds to cellular polynucleotides which

XX enhance and/or are required for viral infectivity, replicative

XX ability or chronicity. The antisense polynucleotides may also be

XX designed to bind with high specificity, to be of increased stability,

XX to be stable and to have low toxicity. The composition also comprises

XX an agent which causes viral RNA to be inactive. The composition

XX is used for preventing HCV replication in a system. The present

XX sequence is encoded by a novel HCV cDNA sequence, which is used in the

XX course of the invention.

XX Sequence 2772 AA;

XX Query Match 99.0%; Score 943; DB 21; Length 2772;

XX Best Local Similarity 98.4%; Prod. No. 1.5e-88;

XX Matches 179; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 MAPITAYAOQTGRLGCGIITSLTGRDNKQVEGEVQIVSTAAQTFLATCINGVCTVYHGA 60

Db 912 LAPITAYAOQTGRLGCGIITSLTGRDNKQVEGEVQIVSTAAQTFLATCINGVCTVYHGA 971

QY 61 GRTITASPQGPVQIOMYTNVDKLVGWPAPQGSRLTPTCTCGSSDLVYVTRHADVIPVRRR 120

Db 972 GRTITASPQGPVQIOMYTNVDKLVGWPAPQGSRLTPTCTCGSSDLVYVTRHADVIPVRRR 1031

QY 121 GDSRGLLSPPRISYLGSGGSPILCPAGHAGVIFRAAVCTRGVAKAVDFIPVESLETTM 180

Db 1032 GDSRGLLSPPRISYLGSGGSPILCPAGHAGVIFRAAVCTRGVAKAVDFIPVESLETTM 1091

QY 181 RS 182

Db 1092 RS 1093

RESULT 13

AAR34009

ID AAR34009 standard; Protein: 2816 AA.

XX AAR34009;

XX 25-MAR-2003 (updated)

XX 26-JUL-1993 (first entry)

XX HCV-1 polypeptide.

XX Polymerase chain reaction; PCR; amplify; primer; hepatitis C virus;

KW HCV; asymptomatic; chronically infected; epitope; viral isolate;
 KW domain; immunological; cross-reactive; envelope protein; vaccine;
 KW gp53(RVDV)/gp55; hog cholera virus; pestivirus; NS5; flavivirus;
 XX

OS Hepatitis C virus.

XX WC9306126-A1.

PN 01-APR-1993.

PF 11-SEP-1992; 92WO-0807683.

PR 13-SEP-1991; 91US-3759575.

PA (CHIR) CHIRON CORP.

PI Houghton M, Weiner AJ;

DR WPI; 1993-117468/14.

XX Immunoreactive hepatitis C virus polypeptide compns. - cont'd.
 PT at least 2 sequences from the first variable domain of distinct
 PT HCV isolates

PS Disclosure; Fig 9; 106pp; English.

XX This sequence represents the entire hepatitis C virus polyprotein.
 CC HCV is a member of the flavivirus family and appears to encode a basic
 CC polypeptide domain ("C") at the N-terminal of the viral polyprotein,
 CC followed by two glycoprotein domains ("E1", "E2/NS1"), upstream of the
 CC nonstructural genes NS2 through NS5. See also AAQ39134-48, AAR35942-
 CC 4008 and AAR3688-89.
 CC (Updated on 25-MAR-2003 to correct PN field.)
 XX

SQ Sequence 2816 AA;

Query Match 99.0%; Score 943; DS 14; Length 2816;
 Best Local Similarity 98.4%; Pred. No. 1.7e-88;

Matches 179; Conservative 3; Mismatches 9; Indels 0; Gaps 0;

QY 1 MAPTAYAQTRGLGCIITSLTGKKNQVEGEVQIVSTAQTFLATCINGCVTVYHGA 60

Db 1026 LAPITAYAQTRGLGCIITSLTGKKNQVEGEVQIVSTAQTFLATCINGCVTVYHGA 1085

QY 61 GRTIASPKGPVICYMTNVDKDLVGNPAPQGSRSITPTCGSSDLYLVTRHADVIPVRRR 120

Db 1086 GRTIASPKGPVICYMTNVDKDLVGNPAPQGSRSITPTCGSSDLYLVTRHADVIPVRRR 1145

QY 121 GDSRGSLSLSPRISYLYKSGSGGPLLCFAGHAGVGFRAAVCTRGVAKAVDFIPVESLETTM 180

Db 1146 GDSRGSLSLSPRISYLYKSGSGGPLLCFAGHAGVGFRAAVCTRGVAKAVDFIPVENLETTM 1205

QY 181 RS 182

Db 1206 RS 1207

RESULT 14

ID AAR70230

XX AAR70230 standard; protein: 2894 AA.

AC AAR70230;

DT 25-MAR-2003 (updated)

DT 07-NOV-1995 (first entry)

DE Composite hepatitis C virus (HC-J1/CDC/CHI).

XX Composite hepatitis C virus; HC-J1/CDC/CHI; HCV; non-A non-B;
 KW synthetic antigens; blood screening.

XX Hepatitis C virus.

OS

XX

PN EP644232-A1.

XX 22-MAR-1995.

XX 14-DEC-1990; 94EP-0108611.

XX 14-DEC-1990; 90EP-0124241.

XX 14-DEC-1990; 90EP-0108611.

XX (INNO-) INNOGENETICS NV.

XX Deleys RJ, Maertens G, Pollet D, Van Heuverswyn H;

XX WPI; 1995-116946/16.

XX Synthetic antigens for the detection of hepatitis C virus
 PT antibodies - comprise portions of the HCV peptide sequence, for
 PT use in screening blood and blood products

PS Disclosure; Fig 1; 51pp; English.

XX AAR70230 is the composite hepatitis C virus (HC-J1/CDC/CHI) protein
 CC from which the synthetic HCV antigens described in AAR70210-R70229
 CC were derived. These synthetic antigens can be used to screen blood,
 CC or blood products for the presence of HCV, they can also be used in
 CC various specific assays for the detection of HCV antibodies, and
 CC antigens, or as immunogens.
 CC (Updated on 25-MAR-2003 to correct PN field.)
 CC (Updated on 25-MAR-2003 to correct PF field.)
 XX

SQ Sequence 2894 AA;

Query Match 99.0%; Score 943; DS 16; Length 2894;
 Best Local Similarity 98.4%; Pred. No. 1.7e-88;

Matches 179; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 MAPTAYAQTRGLGCIITSLTGKKNQVEGEVQIVSTAQTFLATCINGCVTVYHGA 60

Db 1026 LAPITAYAQTRGLGCIITSLTGKKNQVEGEVQIVSTAQTFLATCINGCVTVYHGA 1085

QY 61 GRTIASPKGPVICYMTNVDKDLVGNPAPQGSRSITPTCGSSDLYLVTRHADVIPVRRR 120

Db 1086 GRTIASPKGPVICYMTNVDKDLVGNPAPQGSRSITPTCGSSDLYLVTRHADVIPVRRR 1145

QY 121 GDSRGSLSLSPRISYLYKSGSGGPLLCFAGHAGVGFRAAVCTRGVAKAVDFIPVESLETTM 180

Db 1146 GDSRGSLSLSPRISYLYKSGSGGPLLCFAGHAGVGFRAAVCTRGVAKAVDFIPVENLETTM 1205

QY 181 RS 182

Db 1206 RS 1207

RESULT 15

AAV14975

XX AAV14975 standard; Protein: 2955 AA.

AC AAV14975;

XX 20-MAR-2003 (updated)

XX 08-NOV-1999 (first entry)

XX Amino acid sequence of HCV-1 ORF.

XX Hepatitis C virus; HCV; J1; J7; HCV-1; non-A, non-B HCV; NANBH;
 KW HCV infection; vaccine.

XX Hepatitis C virus.

XX Key Location/Qualifiers

FT Misc-difference 441

FT /note= "encoded by TT"

FT Misc-difference 461

/note- "encoded by CCCC"

FT XX EP339128-A2.
PN XX
XX XX
PD XX 01-SEP-1999.
XX XX 17-SEP-1999: 99EP-0101746.
XX XX 15-SEP-1999: 89US-0498245.
PR XX 21-DEC-1989: 89US-0456142.
PR XX 17-SEP-1999: 90EP-0313149.
XX XX (CHIR) CHIRON CORP.
PA XX (CYAA/) OYA A.
XX XX
PI Cha T, Han J, Houghton M, Irvine RD, Kolberg JA:
PI Miyamura T, Saito I, Weiner AJ:
XX XX
DR WPI: J999-440843/42.
DR N-PSDB: RA207656.
XX XX
PT New Hepatitis C Virus isolates, useful for diagnosis of Hepatitis
PT Infections and development of vaccines
XX XX
PS Disclosure: Fig 12: 142pf: English.
XX XX
CC The invention provides two new isolates of Hepatitis C virus (HCV), H1
CC and J7. These two isolates comprise nucleotide and amino acid sequences
CC that are distinct from the HCV isolate HCV-1. The nucleotide sequences
CC may be used to detect non-A, non-B HCV (NANBH) polynucleotides by
CC hybridisation for diagnosis of NANBH infections. They may also be used to
CC screen blood donors, donated blood and blood products for this infection.
CC The isolates may also be used to isolate other naturally occurring
CC variants of the virus. The polypeptides may be used as a vaccine for
CC administration to patients to protect against infection with NANBH. The
CC present sequence represents the amino acid sequence of HCV-1 ORF.
CC (Updated on 20-MAR-2003 to correct pf field.)
CC (Updated on 20-MAR-2003 to correct PR field.)
XX XX
SQ Sequence 2955 AA;

Query Match 95.0%; Score 943; DB 20; Length 2955;
Best Local Similarity 98.4%; Prod. No. 1.7e-98;
Matches 179; Conservative 3; Mismatches 0; Indels 0; Gaps 0;
QY 1 MAPITAYAOQIRGLLCITITSLTGRDNQVGEVIVSTAAQTFATCINGVGVYHGA 60
Db 1026 LAPITAYAOQIRGLLCITITSLTGRDNQVGEVIVSTAAQTFATCINGVGVYHGA 1085
QY 61 GTRTASPKGFVIOYTVNVDQLVGNPAPOGSRSLTFCGSSDLYLVTHADVIVPVR 120
Db 1086 GTRTASPKGFVIOYTVNVDQLVGNPAPOGSRSLTFCGSSDLYLVTHADVIVPVR 1145
QY 121 GDSRGSLLSPRISYLVKSGSGPLLCFAGHAGVCFRAAVCTRGVAKAVDFIPVESLETTM 180
Db 1146 GDSRGSLLSPRISYLVKSGSGPLLCFAGHAGVCFRAAVCTRGVAKAVDFIPVENLETTM 1205
QY 181 RS 182
Db 1206 RS 1207

Search completed: September 27, 2003, 12:16:53
Job time : 79 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2003 CompuLink Ltd.

OM protein - protein search, using sw model

Run on: September 29, 2003, 18:58:12 : Search time 32 seconds
(without alignments)
44,642 Million cell updates/sec

Title: US-09-965-594-1_COPY_14_22

Perfect score: 44

Sequence: 1 LNC617SL 9

Scoring table: BLASTM62
Gapop 10.0, Gapext 0.5

Searched: 1107863 seqs, 1563726573 residues

Total number of hits satisfying chosen parameters: 1107572

Minimum DB seq length: 0

Maximum DB seq length: 2030000000

Post-processing: Minimum Match 0%

Maximum Match 40%

Listing first 45 summaries

Database: A_Geneseq_19Jun03:

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2: /SIDSL/gcgdata/geneseq/geneseq-emb1/AA1981.DAT:
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8: /SIDSL/gcgdata/geneseq/geneseq-emb1/AA1987.DAT:
9: /SIDSL/gcgdata/geneseq/geneseq-emb1/AA1988.DAT:
10: /SIDSL/gcgdata/geneseq/geneseq-emb1/AA1989.DAT:
11: /SIDSL/gcgdata/geneseq/geneseq-emb1/AA1990.DAT:
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13: /SIDSL/gcgdata/geneseq/geneseq-emb1/AA1992.DAT:
14: /SIDSL/gcgdata/geneseq/geneseq-emb1/AA1993.DAT:
15: /SIDSL/gcgdata/geneseq/geneseq-emb1/AA1994.DAT:
16: /SIDSL/gcgdata/geneseq/geneseq-emb1/AA1995.DAT:
17: /SIDSL/gcgdata/geneseq/geneseq-emb1/AA1996.DAT:
18: /SIDSL/gcgdata/geneseq/geneseq-emb1/AA1997.DAT:
19: /SIDSL/gcgdata/geneseq/geneseq-emb1/AA1998.DAT:
20: /SIDSL/gcgdata/geneseq/geneseq-emb1/AA1999.DAT:
21: /SIDSL/gcgdata/geneseq/geneseq-emb1/AA2000.DAT:
22: /SIDSL/gcgdata/geneseq/geneseq-emb1/AA2001.DAT:
23: /SIDSL/gcgdata/geneseq/geneseq-emb1/AA2002.DAT:
24: /SIDSL/gcgdata/geneseq/geneseq-emb1/AA2003.DAT:
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pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARY

Result No.	Score	Match	Length	DB ID	Description
1	43	97.7	3010	15	AA1953417
2	42	95.5	273	15	AA1950072
3	42	95.5	1051	15	AA1950066
4	42	95.5	1051	16	AA1950061
5	41	93.2	3010	20	AA1950022
6	41	93.2	3010	22	AA1951170
7	41	93.2	3010	22	AA1951174
8	40	90.9	184	21	AA144727
9	40	90.9	631	20	AA1953482

10	39	88.6	3010	14	AA19534580	Human hepatitis C
11	39	88.6	3011	14	AA19534488	Encoded by full-le
12	37	84.1	174	13	AA1950834	HCV NS2-NS4 peptid
13	37	84.1	213	20	AA1951888	HCV NS4A-NS3 compl
14	37	84.1	215	20	AA19517890	HCV NS4A-NS3 compl
15	37	84.1	216	20	AA19517873	HCV NS4A-NS3 compl
16	37	84.1	216	20	AA19517882	HCV NS4A-NS3 compl
17	37	84.1	216	20	AA19517883	HCV NS4A-NS3 compl
18	37	84.1	216	20	AA19517886	HCV NS4A-NS3 compl
19	37	84.1	216	20	AA19517878	HCV NS4A-NS3 compl
20	37	84.1	216	20	AA19517878	HCV NS4A-NS3 compl
21	37	84.1	646	20	AA19524950	HCV NS4A-NS3 compl
22	37	84.1	665	20	AA19524941	HCV NS4A-NS3 compl
23	37	84.1	665	20	AA19524942	HCV NS4A-NS3 compl
24	37	84.1	665	20	AA19524945	HCV NS4A-NS3 compl
25	37	84.1	665	20	AA19524946	HCV NS4A-NS3 compl
26	37	84.1	667	20	AA19517891	HCV NS4A-NS3 compl
27	36	81.8	300	22	AA19527599	Mouse TZR05 amino
28	36	81.8	2894	13	AA19524440	Composite HCV HC-J
29	35	79.5	76	21	AA19534473	Human secreted pro
30	35	79.5	354	21	AA19519987	Arabidopsis thalia
31	35	79.5	354	21	AA19502888	Arabidopsis thalia
32	35	79.5	473	22	AA19523525	Rabbit EST encoded
33	35	79.5	473	23	AA19525529	Human polypeptide
34	34	77.3	144	23	AA19515525	Rifidobacterium lo
35	34	77.3	181	22	AA19536655	Human ORFX protein
36	34	77.3	339	23	AA19527696	Group B Streptococ
37	34	77.3	340	23	AA19527695	Streptococcus poly
38	34	77.3	345	21	AA19525992	Streptococcus poly
39	34	77.3	356	22	AA19521362	Arabidopsis thalia
40	34	77.3	356	22	AA19525686	Novel human diagno
41	34	77.3	1085	22	AA19527847	S cerevisiae apopt
42	34	77.3	1097	22	AA19525216	Novel human diagno
43	33	75.0	82	23	AA1954761	Human albumin diagn
44	33	75.0	82	23	AA1954762	Human albumin diagn
45	33	75.0	82	23	AA1954762	Human albumin diagn

ALIGNMENTS

RESULT 1
AA1953417
ID AA1953417 standard; Protein: 3010 AA.
XX
AC AA1953417:
XX
DT 17-JAN-1995 (first entry)
XX
DE Blood transmissible NANBH virus protein.
XX
KW Polymerase chain reaction; PCR, amplify; primer: non-A, non-B hepatitis;
KW NANBH; virus; blood transmissible; detection; hepatitis virus: KI-PCR;
KW C100 antibody; HCV RNA; NS5 region.
XX
OS Non-A, non-B hepatitis virus.
XX
XX
FH Key Location/Qualifiers
FT Misc-difference 222 /label= His, Arg
FT Misc-difference 226 /label= Cys, Arg
FT Misc-difference 246 /label= Leu, Phe
FT Misc-difference 263 /label= Asp, Asn
FT Misc-difference 291 /label= Phe, Ser
FT Misc-difference 311 /label= Gly, Asp
FT Misc-difference 398 /label= Ser, Arg, Gly
FT Misc-difference 400

FT /label= Thr, Ala
 FT Misc-difference 405
 FT /label= Gln, Pro, Leu
 FT Misc-difference 410
 FT /label= Lys, Arg
 FT Misc-difference 418
 FT /label= Gly, Asp
 FT Misc-difference 430
 FT /label= Asn, Asp
 FT Misc-difference 436
 FT /label= Phe, Leu
 FT Misc-difference 478
 FT /label= Arg, Lys
 FT Misc-difference 759
 FT /label= Leu, Val
 FT Misc-difference 1017
 FT /label= Ser, Asn
 FT Misc-difference 1036
 FT /label= Thr, Ala
 FT Misc-difference 1056
 FT /label= Glu, Asp
 FT Misc-difference 1201
 FT /label= Met, Thr
 FT Misc-difference 1205
 FT /label= Met, Ile
 FT Misc-difference 1255
 FT /label= Asn, Tyr
 FT Misc-difference 1263
 FT /label= Gly, Asp
 FT Misc-difference 1455
 FT /label= Asn, Asp
 FT Misc-difference 1828
 FT /label= Ala, Thr
 FT Misc-difference 1895
 FT /label= Gly, Arg
 FT Misc-difference 1896
 FT /label= Gly, Ile
 FT Misc-difference 2143
 FT /label= Glu, Val
 FT Misc-difference 2144
 FT /label= Asp, Gln
 FT Misc-difference 2462
 FT /label= Cys, Arg
 FT Misc-difference 2486
 FT /label= Val, Met
 FT Misc-difference 2488
 FT /label= Lys, Gln
 FT Misc-difference 2844
 FT /label= Leu, Met
 FT Misc-difference 2862
 FT /label= Leu, Gln
 FT Misc-difference 2917
 FT /label= Arg, Leu
 FT Misc-difference 2968
 FT /label= Ser, Gly
 FT Misc-difference 2985
 FT /label= Cys, Arg
 FT Misc-difference 2990
 FT /label= Tyr, Cys
 XX
 PN JP0610569C-A.
 XX
 PD 19-APR-1994.
 XX
 PF 10-MAR-1992; 92JP-0051885.
 XX
 PR 10-MAR-1992; 92JP-0051885.
 XX
 PA (KAEN/); KAENNO K.
 XX
 DR WPI: 1994-153130/20.
 DR N-PSDB; AAQ63499.
 XX

FT Biochemical-transmissible non-A non-B hepatitis virus DNA - used for
 FT detection of hepatitis virus
 XX
 PS Claim 1; Page 8-20; 22pp; Japanese.
 XX
 CC This sequence is encoded by the genome of a blood transmissible non-A,
 CC non-B hepatitis (NANBH) virus. The cDNA sequence was isolated using the
 CC primers given in AAQ63500-35. The amplified fragments are used in the
 CC detection of hepatitis virus. The target DNA was isolated from serum
 CC of chronically infected NANBH patients who were C100 antibody-positive
 CC and HCV RNA (NS5 region) positive. Reverse transcription-PCR and PCR
 CC were performed on cDNA and the total human NANBH DNA was constructed
 CC from 23 clones.
 XX
 SQ Sequence 3010 AA;
 Query Match 97.7%; Score 43; DB 15; Length 3010;
 Best Local Similarity 88.9%; Pred. No. 65;
 Matches 8; Conservative 1; Mismatches 0; Indels 0; Gaps 0;
 QY 1 LIGCIIITSL 9
 DB 1039 LIGCIVISL 1047
 RESULT 2
 AAR50072
 ID AAR50072 standard; protein; 273 AA.
 XX
 AC AAR50072;
 XX
 DT 24-MAY-1994 (first entry)
 XX
 DE NANBH virus antigenic fragment #4.
 XX
 KW Antigen; structural; non-structural; non A non B hepatitis virus;
 KW NANBH; patient; plasma; diagnosis; detection; carrier; ss.
 XX
 OS Non A Non B hepatitis virus.
 XX
 PN JP06072778-A.
 XX
 PD 15-MAR-1994.
 XX
 PF 21-JUN-1993; 93JP-0156087.
 XX
 PR 10-JUL-1992; 92JP-0207391.
 XX
 PA (KOKU-) KOKUSAI SHIYAKU KK.
 PA (SANW) SANWA KAGAKU KENKYUSHO CO.
 PA (TOFU) TONEN CORP.
 PA (TOKH-) 2H TOKYOTO RINSHO IGAKU SOGO KENKYUSHO.
 XX
 DR WPI: 1994-128677/16.
 DR N-PSDB; AAQ58817.
 XX
 PI Nucleic acid fragment coding non-A non-B hepatitis virus antigen
 PI - useful in diagnosis of NANBH patient and detection of virus
 PI carrier
 XX
 PS Claim 8; Page 18-19; 37pp; Japanese.
 XX
 CC The sequences given in AAR50068 and AAR50070-82 represent antigens of
 CC structural and non-structural regions of non A non B hepatitis virus
 CC (NANBH). The cDNA encoding these sequences were derived from the
 CC plasma of a NANBH patient by recombinant DNA techniques. These
 CC fragments are useful for the diagnosis of NANBH patients and the
 CC detection of NANBH carriers.
 XX
 SQ Sequence 273 AA;
 Query Match 95.5%; Score 42; DB 15; Length 273;
 Best Local Similarity 88.9%; Pred. No. 9.5;

```

Matches      8;  Conservative      1;  Mismatches      0;  Indels      0;  Gaps      0;

QY      1 LIGCIITSL 9
DB      23 LIGCIITSL 31

RESULT 3
AA054066
ID      AAK54066 standard; Protein: 1051 AA.
XX
AC      AAR54066;
DT      14-FEB-1995 (first entry)
DE      Non-A, non-B hepatitis virus gene #4 product.
XX
KW      Non-A, non-B hepatitis virus; NANBHV; Hepatitis C virus; HCV;
KW      core; ENV; NS1; NS2; NS3; antigen; detection.
XX
OS      Hepatitis C virus.
XX
FH      Key      Location/Qualifiers
FT      Region      1..191
FT      Region      /label= core
FT      Region      192..383
FT      Region      /label= ENV
FT      Region      384..810
FT      Region      /label= NS1
FT      Region      811..1051
FT      Region      /label= NS2-NS3
FT      Region      /note= "NS3 N-terminal"
XX
PN      JP06141870-A.
XX
XX      24-MAY-1994.
XX
XX      12-MAR-1992; 92JP-0088140.
XX
XX      12-MAR-1992; 92JP-0088140.
XX
XX      (SANW ) SANWA KAGAKU KENKYUSHO CO.
PA      (TOFU ) TONEN CORP.
PA      (TOKR-) ZH TOKYOYO RINSHO IGAKU SOGO KENKYUSHO.
XX
XX      WPI: 1994-205028/25.
DR      N-PSDB; AAK64068.
XX
XX      DNA coding a Non-A, non-B hepatitis virus antigen - useful for
XX      detecting HCV within serum.
XX
XX      Claim 1-5; Page 11-15; 22pp; Japanese.
XX
XX      Hepatitis C virus #4 and #6 genes were isolated (AAQ64068-69).
CC      Both genes contain the core, ENV, NS1, NS2 and NS3 regions.
CC      A core region fragment is given in AAK64067.
XX
XX      Sequence      1051 AA;

Query Match      95.5%; Score 42; DB 15; Length 1051;
Best Local Similarity      88.9%; Pred. No. 35;
Matches      8;  Conservative      1;  Mismatches      0;  Indels      0;  Gaps      0;

QY      1 LIGCIITSL 9
DB      1039 LIGCIITSL 1047

RESULT 4
AA098361
ID      AAR98361 standard; Protein: 1051 AA.
XX
AC      AAR98361;

```

```

XX      22-AUG-1996 (first entry)
DT
XX
XX      5'UTR/CORE/ENV/NS1/NS2/NS3 from HCV (#4).
DE
XX
XX      Hepatitis C virus; HCV; antigen; detection; antibody.
KW
XX
XX      Hepatitis C virus.
OS
XX
XX      Key      Location/Qualifiers
XX      Peptide      1..191
XX      Peptide      /label= Core peptide
XX      Peptide      192..383
XX      Peptide      /label= ENV1
XX      Peptide      384..810
XX      Peptide      /label= NS1/NS2
XX      Peptide      811..1051
XX      Peptide      /label= NS2 and NS3
XX
XX      JP0713329-A.
XX
XX      23-MAY-1995.
XX
XX      18-JUN-1993; 93JP-0147944.
XX
XX      18-JUN-1993; 93JP-0147944.
XX
XX      (TOFU ) TONEN CORP.
XX
XX      WPI: 1995-228780/25.
XX      N-PSDB; AAT30386.
XX
XX      Recombinant polypeptide comprising partial NS1 region of hepatitis
XX      non-A non-B viral antigen - used in a method for detecting
XX      antibodies against hepatitis non-A non-B virus.
XX
XX      Disclosure; Page 10-12; 15pp; Japanese.
XX
XX      The sequences given in AAR98361-62 represent the 5'UTR/CORE/ENV/NS1/NS2/
XX      NS3 protein region derived from hepatitis C virus (HCV) isolates #4
XX      and #6 respectively. The proteins partic. contain amino acids 384-
XX      495 of the HCV NS1 antigen. These protein fragments may be used in
XX      the detection of antibodies against HCV.
XX
XX      Sequence      1051 AA;

Query Match      95.5%; Score 42; DB 16; Length 1051;
Best Local Similarity      88.9%; Pred. No. 35;
Matches      8;  Conservative      1;  Mismatches      0;  Indels      0;  Gaps      0;

QY      1 LIGCIITSL 9
DB      1039 LIGCIITSL 1047

RESULT 5
AA098022
ID      AAW98022 standard; Protein: 3010 AA.
XX
XX      AAW98022;
AC
XX
XX      21-JUN-1999 (first entry)
DT
XX
XX      Infectious hepatitis C virus genotype 1b strain HC-J4 protein.
DE
XX      HCV; infectious clone; infection; diagnosis; therapy; vaccine;
KW      screening; assay; antiviral; virucide.
XX
XX      Hepatitis C virus.
OS
XX
XX      W05904008-A2.
XX
XX      28-JAN-1999.
PD

```

XX PF 16-JUL-1998; 96W0-0514682.
 XX PR 27-JAN-1998; 98US-0014414.
 XX PR 18-JUL-1997; 97US-0033062.
 XX PA (USSH) US DEPT HEALTH & HUMAN SERVICES.
 XX PT Bukh J, Emerson SJ, Purcell RH, Yanagi M;
 DR WPI; 1999-132252/11.
 DR N-PSDB; AAX24843.
 XX PT New isolated hepatitis C virus nucleic acids - used to develop
 PT products for the diagnosis, prevention and treatment of HCV
 PT infections and for developing screening assays
 XX PS Claim 2; Fig 14G-H; 126pp; English.
 CC This protein is encoded by the infectious hepatitis C virus (HCV)
 CC genotype 1b strain HC-J4 genome (see AAX24833). HC-J4 was obtained
 CC from acute phase plasma of a chimpanzee infected with serum
 CC containing HC-J4/91. The infectious nucleic acid sequence can be
 CC used to produce chimeric genomes (see AAX24833) consisting of the
 CC open reading frames of infectious nucleic acid sequences of other
 CC genotypes (including genotypes 1-6) and subtypes (such as 1b, 2a,
 CC 2b, 2c, 3a, 4a-f, 5a and 6a) of HCV. The invention also relates to
 CC the introduction of mutations or deletions into infectious nucleic
 CC acid sequences in order to produce an attenuated HCV virus suitable
 CC for vaccine development. Infectious nucleic acid sequences can also
 CC be used to produce attenuated virus via passage in vitro or in vivo
 CC of the viruses produced by transfection of a host cell with the
 CC infectious nucleic acid sequence. Vaccines comprising one or more
 CC polypeptides made from the infectious nucleic acid sequence are
 CC used to immunise mammals, especially humans, against hepatitis C.
 CC The nucleic acid sequences can also be used to induce protective
 CC immunity against the virus. The nucleic acid sequences or their
 CC encoded proteases (e.g. NS3 protease) can additionally be used to
 CC develop screening assays to identify antiviral agents for HCV.
 XX SQ Sequence 3010 AA;
 Query Match 93.2%; Score 41; DB 20; Length 3010;
 Best Local Similarity 88.9%; Pred. No. 1.5e+02;
 Matches 8; Conservative 1; Mismatches 0; Indels 0; Gaps 0;
 QY 1 LAGCIITSL 9
 DB 1039 VLGCIIITSL 1047
 RESULT 6
 AAB31170
 ID AAB31170 standard; Protein; 3010 AA.
 XX AC AAB31170;
 XX DT 02-APR-2001 (first entry)
 XX DE Amino acid sequence of a hepatitis C virus (HCV) clone genotype 1b.
 XX KW Chimeric virus; bovine viral diarrhoea virus; BVDV; hepatitis C virus;
 XX KW HCV; vaccine; viral inhibitor; antiviral.
 XX OS Hepatitis C virus.
 XX PN WO200075352-A2.
 XX PD 14-DEC-2000.
 XX PF 02-JUN-2000; 2000W0-US15527.
 XX PR 04-JUN-1999; 99US-0137817.

XX PA (USSH) US DEPT HEALTH & HUMAN SERVICES.
 XX PT Nam J, Bukh J, Emerson SJ, Purcell RH;
 DR WPI; 2001-071081/08.
 DR N-PSDB; AAC86939.
 XX PT New nucleic acid comprising a chimeric bovine viral diarrhoea virus
 PT genome in which the (non-)structural region has been replaced by
 PT hepatitis C virus (HCV) genome useful for treating or preventing HCV
 PT signs and symptoms
 XX PS Disclosure; Fig 4G-H; 97pp; English.
 CC The specification describes a nucleic acid comprising a chimeric virus
 CC genome, specifically bovine viral diarrhoea virus (BVDV) genome in which
 CC the (non-)structural region has been replaced by the (non-)structural
 CC region of a hepatitis C virus (HCV) genome. The nucleic acids comprising
 CC the chimeric virus and the chimeric virus are useful for identifying
 CC cell lines capable of supporting the replication of these chimeric
 CC viruses, in screening for neutralizing antibodies to HCV of different
 CC genotypes, in the production of HCV-BVDV virions, for the development
 CC of inactivated or attenuated vaccines to prevent HCV-BVDV in a mammal,
 CC in studying the molecular properties of HCV indirectly in vitro, and in
 CC identifying inhibitors of viral enzyme activity which would be useful
 CC as antiviral agents. Formulations or compositions comprising the
 CC chimeric virions may be used to treat or prevent the signs and symptoms
 CC of HCV. The present sequence is encoded by a HCV clone, which is used
 CC to construct chimeric nucleic acids of the invention.
 XX SQ Sequence 3010 AA;
 Query Match 93.2%; Score 41; DB 22; Length 3010;
 Best Local Similarity 88.9%; Pred. No. 1.5e+02;
 Matches 8; Conservative 1; Mismatches 0; Indels 0; Gaps 0;
 QY 1 LAGCIITSL 9
 DB 1039 VLGCIIITSL 1047
 RESULT 7
 AAB59174
 ID AAB59174 standard; Protein; 3010 AA.
 XX AC AAB59174;
 XX DT 21-MAR-2001 (first entry)
 XX DE Protein encoded by infectious Hepatitis C virus 1b genotype.
 XX KW BVDV-B; hepatitis C virus; HCV; vaccine.
 XX OS Hepatitis C virus.
 XX PN WO200075337-A1.
 XX PD 14-DEC-2000.
 XX PF 02-JUN-2000; 2000W0-US15293.
 XX PR 04-JUN-1999; 99US-0137564.
 XX PA (USSH) US DEPT HEALTH & HUMAN SERVICES.
 XX PT Bukh J, Yanagi M, Emerson SJ, Purcell RH;
 DR WPI; 2001-091214/10.
 XX PT New infectious nucleic acids of the GB virus-B clone, useful for
 PT indirectly studying the molecular properties of hepatitis C virus (HCV)
 PT and in developing vaccines and therapeutics for HCV

XX PS Disclosure; Fig 7; 96pp; English.

XX CC The present invention relates to GB virus-B. The nucleic acid molecules of the invention are useful for indirectly studying the molecular properties of hepatitis C virus (HCV). The infectious nucleic acid sequence of the GB virus-B clone and the HCV/GBV-B chimera may be used in the development of vaccines and therapeutics for HCV.

XX CC Sequence 3010 AA:

Query Match 93.2%; Score 41; DB 22; Length 3010;
Best Local Similarity 88.5%; Pred. No. 1.5e-02;
Matches 8; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

OY 1 LGCIIITSL 9
:|||||

DB 1039 VLGCIIITSL 1047

RESULT 8
AA44727
ID AAY44727 standard; Protein: 184 AA.
AC AAY44727;
DI 04-MAY-2000 (first entry)
XX Hepatitis C virus NS3 catalytic domain of 1B genotype.
XX NS3 catalytic domain; NS4A peptide; NS4A-NS3 fusion construct; diagnosis;
KW serine protease; trypsin family; screening; anti-viral compound;
KW treatment; inhibitor; therapeutic.
XX Hepatitis C virus.
XX Key Location/Qualifiers
FH 1..5
FT Region
FT /note= "Residues that are deleted for the construction
FT of NS4A/NS3 fusion construct"

XX PN WO200001718-A2.
XX 13-JAN-2000.
XX 02-JUL-1999; 98WO-US15035.
XX 02-JUL-1998; 98US-0091675.
XX (UYFL) UNIV FLORIDA.
XX Dunn BM, Bukhtiyarova M;
XX WPI; 2000-182103/16.
XX Novel polypeptide comprising hepatitis C virus NS4A and NS3 domains.
XX useful for screening for compounds useful for the diagnosis and
XX treatment of hepatitis C virus.

XX PS Claim 3; Fig 1; 30pp; English.

XX CC The present protein sequence is the catalytic domain of the NS3 clone
XX from the 1B genotype, derived from hepatitis C virus. The NS3 protein
XX has a sequence relationship to members of the trypsin family of serine
XX proteases. Constructs comprising an NS4A peptide sequence covalently
XX attached to the N-terminus of NS3 catalytic domain or their fragments
XX can be used to create fusion polypeptides. This fusion polypeptide
XX contains the NS3 domain expressed in a stable, soluble form. This
XX facilitates the use of the polypeptide in direct screening of potential
XX anti-viral compounds, that are used for diagnosis and treatment of
XX hepatitis C virus infection. It is also used to screen for inhibitors of
XX serine protease activity. The polynucleotides are also useful to identify
XX diagnostic or therapeutic compounds and for recombinant production of

XX CC the fusion polypeptide.

XX SO Sequence 184 AA;

Query Match 90.9%; Score 40; DB 21; Length 184;
Best Local Similarity 100.0%; Pred. No. 15;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 2 LGCIIITSL 9
:|||||

DB 15 LGCIIITSL 22

RESULT 9
AAK3482
ID AAK93482 standard; Protein: 631 AA.
AC AAK93482;
XX 11-JUN-1999 (first entry)
XX HCV NS3 protein.
XX NS3; helicase domain; X-ray crystal structure; Hepatitis C virus;
KW HCV; computer programme; binding pocket; three-dimensional.
XX Hepatitis C virus.
XX WO9909148-A1.
XX 25-FEB-1999.
XX 13-AUG-1998; 98WO-US16879.
XX 13-AUG-1997; 97US-0055772.
XX (VERT-) VERTEX PHARM INC.
XX Caron P, Kim J, Lin C, Morgenstern K;
XX WPI; 1999-190157/16.
XX N-PSDB; AAK21258.
XX New hepatitis C virus NS3 helicase crystals - which provide
XX molecular design techniques to identify, select and design agents
XX which bind to the helicase, particularly inhibitor compounds
XX Disclosure; Page 202-205; 224pp; English.
XX This invention relates to the X-ray crystal structure of the Hepatitis C
XX virus helicase domain. The invention describes crystallized complexes of
XX HCV helicase and an oligonucleotide. The described method is used in a
XX novel computer programme where the computer is programmed with the
XX structure coordinates of the HCV helicase oligonucleotide binding pocket
XX or the HCV helicase nucleotide triphosphate pocket wherein the computer
XX is capable of displaying a three-dimensional representation of that
XX binding pocket.

XX SO Sequence 631 AA;

Query Match 90.9%; Score 40; DB 20; Length 631;
Best Local Similarity 100.0%; Pred. No. 49;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 2 LGCIIITSL 9
:|||||

DB 14 LGCIIITSL 21

RESULT 10
AAK34580
ID AAK34580 standard; Protein: 3010 AA.
XX

CC were used in the isolation of the NS2-NS4 regions of the Hepatitis C
 CC Virus (HCV) gene of the invention (see also: AAR29660, AAR29559-60 and
 CC AAR29843-51). These RNA sequences were isolated from the serum of a
 CC patient suffering from hepatitis C (HC). The isolated RNA sequences
 CC were converted into cDNA using transcriptase in the presence of one
 CC of the primer sequences given in AAG32578-79. The sequences were then
 CC amplified using primer pairs. The cDNA sequences isolated represent
 CC different alleles of the same region of the HCV gene. Sequence
 CC comparisons of these clones showed that it is possible for a patient
 CC to carry more than one HCV strain at one time. See also AAG32436.
 CC (Updated on 25-MAR-2003 to correct PN field.)
 XX
 XX

SQ Sequence 174 AA:

Query Match 84.1% Score 37; DB 13; Length 174;
 Best Local Similarity 88.9%; Pred. No. 49;
 Matches 8; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Oy 1 ILGCKITSL 9
 Db 55 LLGCKITSL 64
 --- Not a match

RESULT 13
 AAY17889
 ID AAY17886 standard; Protein: 215 AA.

XX AC AAY17889;

XX DT 07-SEP-1999 (first entry)

XX DE HCV NS4A-NS3 complex SEQ ID NO:50.

XX KW HCV; hepatitis C virus; single chain recombinant complex; linker;
 KW NS3 protease; NS4A cofactor; NS4A-NS3 complex; serine protease;
 KW hydrophobic domain; covalent complex; detection; inhibitor.

OS Hepatitis C virus.

OS Synthetic.

XX PN W09928482-A2.

XX PD 10-JUN-1999.

XX PF 24-NOV-1998; 98WO-US24528.

XX PR 28-JUL-1998; 98US-0094331.

XX PR 28-NOV-1997; 97US-0067315.

XX PA (SCHE) SCHERING CORP.

PI Malcolm BA, Taremi SS, Weber PC, Yao N;

XX WPI; 1999-385385/32.

XX New hepatitis C virus covalent complexes

XX Example 1; Page 119; 21pp; English.

CC The present invention describes a covalent hepatitis C virus (HCV)
 CC NS4A-NS3 complex comprising a central hydrophobic domain of native HCV
 CC NS4A peptide, a linker, and an HCV NS3 serine protease domain, where the
 CC hydrophobic domain of native HCV NS4A peptide is tethered by the linker
 CC to the amino terminus of the HCV NS3 protease domain. The present
 CC sequence represents an example of the above complex. The covalent
 CC NS4A-NS3 complexes are useful for structural determination and
 CC determination of mode of binding of HCV inhibitors by NMR spectroscopy.
 CC They can also be used for detecting inhibitors of the protease activity,
 CC the helicase activity and the ATPase activity of NS3. The covalent
 CC NS4A-NS3 complexes are more soluble, stable and active than the non
 CC covalent protease-peptide complexes previously available.

XX Sequence 213 AA:

Query Match 84.1% Score 37; DB 20; Length 213;
 Best Local Similarity 88.9%; Pred. No. 60;
 Matches 8; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Oy 1 ILGCKITSL 9
 Db 47 LLGCKITSL 55

RESULT 14
 AAY17890
 ID AAY17890 standard; Protein: 215 AA.

XX AC AAY17890;

XX DT 07-SEP-1999 (first entry)

XX DE HCV NS4A-NS3 complex SEQ ID NO:57.

XX KW HCV; hepatitis C virus; single chain recombinant complex; linker;
 KW NS3 protease; NS4A cofactor; NS4A-NS3 complex; serine protease;
 KW hydrophobic domain; covalent complex; detection; inhibitor.

OS Hepatitis C virus.

OS Synthetic.

XX PN W09928482-A2.

XX PD 10-JUN-1999.

XX PF 24-NOV-1998; 98WO-US24528.

XX PR 28-JUL-1998; 98US-0094331.

XX PR 28-NOV-1997; 97US-0067315.

XX PA (SCHE) SCHERING CORP.

PI Malcolm BA, Taremi SS, Weber PC, Yao N;

XX WPI; 1999-385385/32.

XX New hepatitis C virus covalent complexes

XX Example 1; Page 122-123; 21pp; English.

CC The present invention describes a covalent hepatitis C virus (HCV)
 CC NS4A-NS3 complex comprising a central hydrophobic domain of native HCV
 CC NS4A peptide, a linker, and an HCV NS3 serine protease domain, where the
 CC hydrophobic domain of native HCV NS4A peptide is tethered by the linker
 CC to the amino terminus of the HCV NS3 protease domain. The present
 CC sequence represents an example of the above complex. The covalent
 CC NS4A-NS3 complexes are useful for structural determination and
 CC determination of mode of binding of HCV inhibitors by NMR spectroscopy.
 CC They can also be used for detecting inhibitors of the protease activity,
 CC the helicase activity and the ATPase activity of NS3. The covalent
 CC NS4A-NS3 complexes are more soluble, stable and active than the non-
 CC covalent protease-peptide complexes previously available.

SQ Sequence 215 AA:

Query Match 84.1% Score 37; DB 20; Length 215;
 Best Local Similarity 88.9%; Pred. No. 60;
 Matches 8; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Oy 1 ILGCKITSL 9
 Db 47 LLGCKITSL 55

RESULT 15
 AAY17879
 ID AAY17879 standard; Protein: 216 AA.

```

XX  AAV17879;
AC
XX  07-SEP-1999 (first entry)
XX
XX  HCV NS4A-NS3 complex SEQ ID NO:3.
XX
XX  HCV: hepatitis C virus; single chain recombinant complex; linker:
XX  NS3 protease; NS4A cofactor; NS4A-NS3 complex; serine protease;
XX  hydrophobic domain; covalent complex; detection; inhibitor.
XX
XX  Hepatitis C virus.
XX  Synthetic.
XX
XX  WO9928482-A2.
XX
XX  10-JUN-1999.
XX
XX  24-NOV-1998; 98WO-0524528.
XX
XX  28-JUL-1998; 98US-0034331.
XX  28-NOV-1997; 97US-0067315.
XX
XX  (SCHE ) SCHERING CORP.
XX
XX  Malcolm BA, Faremi SS, Weber PC, Yao N;
XX
XX  WP: 1999-385385/32.
XX
XX  New hepatitis C virus covalent complexes
XX
XX  Claim 6: Page 75-76; 21pp; English.
XX
XX  The present invention describes a covalent hepatitis C virus (HCV)
XX  NS4A-NS3 complex comprising a central hydrophobic domain of native HCV
XX  NS4A peptide, a linker, and an HCV NS3 serine protease domain, where the
XX  hydrophobic domain of native HCV NS4A peptide is tethered by the linker
XX  to the amino terminus of the HCV NS3 protease domain. The present
XX  sequence represents a specifically claimed example of the above
XX  complex. The covalent NS4A-NS3 complexes are useful for structural
XX  determination and determination of mode of binding of HCV inhibitors by
XX  NMR spectroscopy. They can also be used for detecting inhibitors of the
XX  protease activity, the helicase activity and the ATPase activity of NS3.
XX  The covalent NS4A-NS3 complexes are more soluble, stable and active than
XX  the non-covalent protease-peptide complexes previously available.
XX
XX  Sequence 216 AA:
XX
XX  Query Match 84.1% Score 37; DB 20; Length 216;
XX  Best Local Similarity 88.9%; Pred. No. 61;
XX  Matches 8; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
XX
XX  QY 1 ILSCIIITSL 9
XX  11111111
XX  Db 48 LGGCKTSL 56
XX
XX  Search completed: September 29, 2003, 19:07:02;
XX  Job time : 33 secs

```

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: September 29, 2003, 19:03:22 : Search time 13 seconds
(without alignments)
48.084 Million cell updates/sec

Title: US-09-965-594-1_COPY_14_22

Perfect score: 44
Sequence: 1 LLGCIITSL 9

Scoring table: BLOSUM62
Gapop 10.0, Gapext 0.5

Searched: 29330w seqs, 5616062 residues

Total number of hits satisfying chosen parameters: 293303

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%

Maximum Match 96%

Listing first 45 summaries

Database: PIR761*

1: pir1*

2: pir2*

3: pir3*

4: pir4*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Query	Score	Match	Length	DB ID	Description
1	43	97.7	3010	1	A45573	genome polyprotein
2	40	90.9	3010	1	GNMW1W	genome polyprotein
3	39	88.6	3010	1	S18030	genome polyprotein
4	38	86.4	231	2	I16938	hypothetical prote
5	37	84.1	851	2	G71435	hypothetical prote
6	36	81.8	403	2	B69338	conserved hypotet
7	35	79.5	653	2	A49722	endoglin precursor
8	34	77.3	179	2	A01917	hypothetical prote
9	34	77.3	338	2	H86372	protein F508.40.1
10	34	77.3	345	2	G86372	protein F508.39.1
11	34	77.3	428	2	PC4163	toxin-co-regulated
12	34	77.3	946	2	S48255	probable membrane
13	34	77.3	2783	2	T31431	inositol 1,4,5-tri
14	33	75.0	350	1	S00755	pleckstrin - human
15	33	75.0	397	2	D83311	conserved hypotet
16	33	75.0	429	2	A61463	flagellar biosynth
17	33	75.0	441	2	A69744	beta-lactamase hom
18	33	75.0	472	2	G71503	probable replicati
19	33	75.0	525	2	T34544	hypothetical prote
20	33	75.0	1056	2	T02930	lysine-ketoglutarate
21	34	75.0	1089	1	PF0064	platelet-derived g
22	32	72.7	158	2	H90459	hypothetical prote
23	32	72.7	258	2	A00459	Sec-independent pr
24	32	72.7	301	1	S12864	retinal isomerase
25	32	72.7	323	2	C89045	protein AC219.5.1
26	32	72.7	330	2	C69648	2-keto-3-deoxygluc
27	32	72.7	348	2	C81994	aldose 1-epimerase
28	32	72.7	367	2	C82943	ferri-chrome ABC tr
29	32	72.7	393	2	D65238	hypothetical_44.4

30	32	72.7	399	2	A60088	postsynaptic membr
31	32	72.7	417	2	A82984	acetyl-CoA C-acety
32	32	72.7	417	2	E98299	probable acyl-CoA
33	32	72.7	419	2	JC4522	alpha-galactosidas
34	32	72.7	438	2	AD0481	probable glucanase
35	32	72.7	442	2	D90122	putative helicase
36	32	72.7	444	2	PD0001	protein-glutamine
37	32	72.7	501	2	T45955	hypothetical prote
38	32	72.7	519	2	T43333	hexose transporter p
39	32	72.7	531	2	T40480	hectose transporter
40	32	72.7	573	2	H86313	protein F2H15.10.1
41	32	72.7	620	2	T13460	hypothetical prote
42	32	72.7	691	1	A29996	protein-glutamine
43	32	72.7	928	2	T36419	Sec76 domain prote
44	32	72.7	1064	2	T05195	saccharopine dehyd
45	32	72.7	1074	2	UC5928	semaphorin F precu

ALIGNMENTS

RESULT 1

A45573
genome polyprotein - hepatitis C virus (strain J7)
N:Contains: capsid protein C; envelope protein M; hepatitis virus (EC 3.4.21.98) (nonst
protein NS4a; nonstructural protein NS4b; nonstructural protein NS5
C:Species: hepatitis C virus
C:Date: 19-May-2000 #sequence_revision: 19-May-2000 #text_change: 19-Jan-2001
C:Accession: A45573
R:Tanaka, T.; Kato, N.; Nakagawa, M.; Ootsuyama, Y.; Cho, M.J.; Nakazawa, T.; Hijik
Virus Res. 23, 39-53, 1992
A:Title: Molecular cloning of hepatitis C virus genome from a single Japanese carri
A:Reference numbers: A45573; MUID:92295714; PMID:1318627
A:Accession: A45573
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-3010 <FAN>
A:Cross-references: GB:D11168; GB:D01171; NID:9221612; PIGN:BAA01943.1; PID:922161
A:Experimental source: HCV-JT
A:Note: sequence extracted from NCBI backbone (NCBI:106206, NCBI:P:106207)
C:Superfamily: hepatitis C virus genome polyprotein
C:Keywords: ATP; glycoprotein; hydrolase; nucleotide binding; P-loop; polyprotein;
P:2 115/Product: capsid protein C #status predicted <CPC>
P:116-191/Product: envelope protein M #status predicted <EPM>
P:192-389/Product: major envelope protein E #status predicted <MEE>
P:390-729/Product: nonstructural protein NS1 #status predicted <NS1>
P:730-1006/Product: nonstructural protein NS2 #status predicted <NS2>
P:1007-1615/Product: hepatitis virus NS2 #status predicted <NS2>
P:1230-1237/Region: nucleotide-binding motif A (P-loop)
P:1312-1317/Region: nucleotide-binding motif A (P-loop)
P:1316-1319/Region: nucleotide-binding motif B
P:1616-1862/Product: nonstructural protein NS4a #status predicted <NS4a>
P:1863-2013/Product: nonstructural protein NS4b #status predicted <NS4b>
P:2014-3010/Product: nonstructural protein NS5 #status predicted <NS5>

Query Match 97.7% Score 43: DB 1: Length 3010;

Best Local Similarity 88.9% Preq. No. 11;

Matches 8; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

Qy 1 LLGCIITSL 9

Db 1039 LLGCIITSL 1047

RESULT 2

GNMW1W

genome polyprotein - hepatitis C virus (strain Taiwan)
N:Contains: capsid protein C; envelope protein M; hepatitis virus (EC 3.4.21.98) (nonst
protein NS4a; nonstructural protein NS4b; nonstructural protein NS5
C:Species: hepatitis C virus
A:Note: host Homo sapiens (man)
C:Date: 31-Dec-1992 #sequence_revision: 31-Dec-1992 #text_change: 19-Jan-2001
C:Accession: A40244

R:Chen, P.-J.; Lin, M.-H.; Tai, K.-F.; Liu, P.-C.; Lin, C.-J.; Chen, D.-S.; Virology 188, 102-113, 1992
 A:Title: The Taiwanese hepatitis C virus genome: sequence determination and mapping the
 A:Reference number: A40244; MUID:92230206; PMID:1214443
 A:Accession: A40244
 A:Molecule type: genomic RNA
 A:Residues: 1-3010 <CH>
 A:Cross-references: GR:M84754
 C:Superfamily: Hepatitis C virus genome polyprotein
 C:Keywords: ATP; capsid protein; envelope protein; glycoprotein; hydrolase; nonstructural
 F:1115/Product: capsid protein C #status predicted <CP>
 F:116-191/Product: envelope protein E #status predicted <EP>
 F:192-389/Product: major envelope protein NS1 #status predicted <NE>
 F:390-729/Product: nonstructural protein NS2 #status predicted <NS2>
 F:730-1006/Product: nonstructural protein NS3 #status predicted <NS3>
 F:1007-1615/Product: nonstructural protein NS4 #status predicted <NS4>
 F:1230-1237/Region: nucleotide-binding motif A (P-loop)
 F:1230-1237/Region: nucleotide-binding motif B
 F:1312-1317/Region: DEXH motif
 F:1316-1319/Region: DEXH motif
 F:1616-1862/Product: nonstructural protein NS4a #status predicted <NS4>
 F:1863-2013/Product: nonstructural protein NS4b #status predicted <NS4b>
 F:2014-3010/Product: nonstructural protein NS5 #status predicted <NS5>
 F:196,209,233,234,250,305,325,417,423,430,448,532,540,556,576,623,645,1253,1255,2041,2073
 Query Match 90.9%; Score 40; DB 1; Length 3010;
 Best Local Similarity 88.9%; Pred. No. 40;
 Matches 8; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 QY 1 LLGCIITSL 9
 DB 1039 LF8CIIITSL 1047
 |||||
 RESULT 3
 S18030
 genome polyprotein - hepatitis C virus (isolate JKI)
 N:Contents: capsid protein C; envelope protein E; hepatitis C virus (nonstructural
 protein NS4a; nonstructural protein NS4b; nonstructural protein NS5
 C:Species: hepatitis C virus
 A:Variety: isolate JKI
 C:Date: 19-May-2000 #sequence_revision 19-May-2000 #text_change 23-Mar-2001
 C:Accession: S18030; S33570; A48332; S18029
 R:Honda, M.; Kaneko, S.; Masahito, U.; Kobayashi, K.; Murakami, S.
 Submitted to the EMBL Data Library, September 1999
 A:Description: A whole genome of hepatitis C virus cDNA was isolated from a single patie
 A:Reference number: S18028
 A:Molecule type: genomic RNA
 A:Residues: 1-3010 <CH>
 A:Cross-references: EMBL:X51596; NID:659478; PDB:CA44733.1; PID:954474
 A:Experimental source: isolate JKI from an individual
 R:Honda, M.; Kaneko, S.; Masahito, U.; Kobayashi, K.; Murakami, S.
 Arch. Virol. 128, 163-169, 1993
 A:Title: Sequence analysis of putative structural regions of hepatitis C virus isolated
 A:Reference number: A48332; MUID:93119270; PMID:8380422
 A:Accession: S33570
 A:Molecule type: genomic RNA
 A:Residues: 1-547, 1-549-621, 1-623-624, 1-626-652, 1-655-761, 1-763-782 <CH>
 A:Cross-references: EMBL:X51591
 A:Note: this sequence is inconsistent with the nucleotide translation
 A:Note: the authors translated the codon ACG for residue 43 as Pro, TAG for residue 120
 as Trp, and TTC for residue 771 as Ser
 A:Note: sequence extracted from NCBI backbone (NCBI:121747, NCBI:121748)
 C:Superfamily: hepatitis C virus genome polyprotein
 C:Keywords: ATP; glycoprotein; hydrolase; nucleotide binding; P-loop; polyprotein; serin
 F:2-115/Product: capsid protein C #status predicted <CP>
 F:116-191/Product: envelope protein E #status predicted <EP>
 F:192-389/Product: major envelope protein E #status predicted <NE>
 F:390-729/Product: nonstructural protein NS1 #status predicted <NS1>
 F:730-1006/Product: nonstructural protein NS2 #status predicted <NS2>
 F:1007-1615/Product: nonstructural protein NS3 #status predicted <NS3>
 F:1230-1237/Region: nucleotide-binding motif A (P-loop)
 F:1312-1317/Region: nucleotide-binding motif B

F:1316-1319/Region: DEXH motif
 F:1616-1862/Product: nonstructural protein NS4a #status predicted <NS4>
 F:1863-2013/Product: nonstructural protein NS4b #status predicted <NS4b>
 F:2014-3010/Product: nonstructural protein NS5 #status predicted <NS5>
 F:196,209,234,250,305,417,423,448,532,540,556,576,623,645/Binding site: carbohydrate
 Query Match 88.6%; Score 39; DB 1; Length 3010;
 Best Local Similarity 77.8%; Pred. No. 61;
 Matches 7; Conservative 1; Mismatches 1; Indels 0; Gaps 0;
 QY 1 LLGCIITSL 9
 DB 1039 LF8CIIITSL 1047
 |||||
 RESULT 4
 T16936
 hypotrichal protein T25D10.2 - Caenorhabditis elegans
 C:Species: Caenorhabditis elegans
 C:Date: 20-Sep-1999 #sequence_revision 20-Sep-1999 #text_change 20-Sep-1999
 C:Accession: T16938
 R:Miller, N.
 Submitted to the EMBL Data Library, June 1995
 A:Description: The sequence of C. elegans cosmid T25D10.
 A:Reference number: Z18611
 A:Accession: T16938
 A>Status: preliminary; translated from GB/EMBL/DBJ
 A:Molecule type: DNA
 A:Residues: 1-231 <ML>
 A:Cross-references: EMBL:U284C9; NID:9849230; PID:9849234; PDB:1AC46592.1; CESP:T2
 A:Experimental source: strain Bristol N2
 C:Genetics:
 A:Gene: CESP:T25D10.2
 A:Introns: 35/3; 56/1; 61/3; 76/3; 105/1; 111/3; 171/1; 201/1
 Query Match 86.4%; Score 38; DB 2; Length 231;
 Best Local Similarity 87.5%; Pred. No. 11;
 Matches 7; Conservative 1; Mismatches 0; Indels 0; Gaps 0;
 QY 1 LLGCIITS 8
 DB 193 LLGCIITS 200
 |||||
 RESULT 5
 G71435
 hypotrichal protein - Arabidopsis thaliana
 C:Species: Arabidopsis thaliana (mouse-ear cress)
 A:Variety: Columbia
 C:Date: 03-Aug-1998 #sequence_revision 03-Aug-1998 #text_change 05-Dec-1998
 C:Accession: G71435
 R:Bevan, M.; Bancroft, T.; Reil, E.; Love, K.; Goodman, H.; Dean, C.; Bergkamp, R.;
 P.; Wedler, H.; Wedler, F.; Wambutt, R.; Weitzenecker, T.; Pohl, T.M.; Terry, N.;
 Vranagh, T.; Hempel, S.; Kottler, P.; Entian, K.D.; Rieger, M.; Schaeffer, M.; Funk,
 Nature 351, 485-488, 1998
 A:Authors: Mueller-Auer, S.; Silvey, M.; James, R.; Montfort, A.; Pons, A.; Puigdom
 erhoft, A.; Moores, T.; Jones, J.D.S.; Eneva, T.; Palmer, K.; Benes, V.; Rechman, S.;
 C.; Chalvatius, N.
 A:Title: Analysis of 1.9 Mb of contiguous sequence from chromosome 4 of Arabidopsis
 A:Reference number: A71400; MUID:98121113; PMID:9461215
 A:Accession: G71435
 A>Status: preliminary; nucleic acid sequence not shown; translation not shown
 A:Molecule type: DNA
 A:Residues: 1-851 <BE>
 A:Cross-references: GB:297342; NID:g2245031; PID:e327517; PID:g2245038
 C:Genetics:
 A:Map position: 4COP9-4C3445
 Query Match 84.1%; Score 37; DB 2; Length 851;
 Best Local Similarity 77.8%; Pred. No. 50;
 Matches 7; Conservative 1; Mismatches 1; Indels 0; Gaps 0;
 QY 1 LLGCIITSL 9

Db 291 LGCTITSL 293
|||||
RESULT 6
B69338
conserved hypothetical protein AFC706 - Archaeoglobus fulgidus
C:Species: Archaeoglobus fulgidus
C>Date: 05-Dec-1997 #sequence_revision 05-Dec-1997 #text_change 22-Oct-1999
C:Accession: B69338
R:Klenk, H.P.; Clayton, R.A.; Tomb, C.F.; White, C.; Nelson, K.E.; Ketchum, K.A.; Dodson
J.; Fleischmann, R.D.; Quackenbush, J.; Lee, N.H.; Sutton, G.G.; Gill, S.; Kirkness, E.F.
Glocke, A.; Zhou, L.; Overbeek, R.; Gocayne, J.D.; Weidman, J.F.; McDonald, L.
Nature 390, 364-370, 1997
A:Authors: Overbeek, R.; Cotton, M.D.; Springs, T.; Atlicha, P.; Kaine, B.P.; Sykes, S.
Smith, H.O.; Woese, C.R.; Venter, J.C.
A:Title: The complete genome sequence of the hyperthermophilic, sulfate-reducing archae
A:Reference number: A69250; MUID:94249343; PMID:9389475
A:Accession: B69338
A>Status: preliminary; nucleic acid sequence not shown; translation not shown
A:Molecule type: DNA
A:Residues: 1-403 <KL>
A:Cross-references: GB:AE001056; GR:AE000722; NID:94562379; PIDN:AA800575; PID:0254991
Query Match 81.8% Score 34; DB 2; Length 403;
Best Local Similarity 66.7% Pred. No. 41;
Matches 6; Conservative 2; Mismatches 1; Indels 0; Gaps 0;
QY 1 LGCTITSL 9
| | | | |
Db 330 IFGCIVTAL 338
|||
RESULT 7
A49722
endoglin precursor - pig
C:Species: Sus scrofa domestica (domestic pig)
C>Date: 03-May-1994 #sequence_revision 03-May-1994 #text_change 05-Nov-1999
C:Accession: A49722; S40180
R:Yamashita, H.; Ichijo, H.; Grimsby, S.; Moran, A.; ten Dijke, P.; Miyazawa, K.
J. Biol. Chem. 269, 1995-2003, 1994
A:Title: Endoglin forms a heteromeric complex with the signaling receptors for transform
A:Reference number: A49722; MUID:94124550; PMID:8294451
A:Accession: A49722
A>Status: preliminary
A:Molecule type: mRNA
A:Residues: 1-553 <YAM>
A:Cross-references: GR:723142; NID:9437919; PIDN:CA86673; PID:0437920
C:Keywords: homodimer; phosphoprotein; signal transduction; transmembrane protein
Query Match 79.5% Score 35; DB 2; Length 653;
Best Local Similarity 87.5% Pred. No. 54;
Matches 7; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 2 LGCTITSL 9
||| ||||
Db 379 LGCTITSL 385
||| ||||
RESULT 8
AD1917
hypothetical protein all0887 [imported] - Nostoc sp. (strain PCC 7120)
C:Species: Nostoc sp. PCC 7120
A:Note: Nostoc sp. strain PCC 7120 is a synonym of Anabaena sp. strain PCC 7120
C>Date: 14-Dec-2001 #sequence_revision 14-Dec-2001 #text_change 09-Dec-2002
C:Accession: AD1917
R:Kaneko, T.; Nakamura, Y.; Wolk, C.P.; Kuri, T.; Sasamoto, S.; Watanabe, A.; Itiguchi
Nakazaki, N.; Shimpo, S.; Sugimoto, M.; Takazawa, Y.; Yamada, M.; Yasuda, M.; Tabata, S.
DNA Res. 8, 205-213, 2001
A:Title: Complete Genomic Sequence of the Filamentous Nitrogen-fixing Cyanobacterium Ana
A:Reference number: AB-807; MUID:21595285; PMID:11759640
A:Accession: AD1917
A>Status: preliminary
A:Cross-references: GB:AE001056; GR:AE000722; NID:94562379; PIDN:AA800575; PID:0254991

A:Molecule type: DNA
A:Residues: 1-179 <KUR>
A:Cross-references: GB:BA000019; PIDN:BAB72844.1; PID:917130232; GSPDB:GN00179
A:Experimental source: strain PCC 7120
C:Genetics:
A:Gene: all0887
Query Match 77.3% Score 34; DB 2; Length 179;
Best Local Similarity 85.7% Pred. No. 49;
Matches 6; Conservative 1; Mismatches 0; Indels 0; Gaps 0;
QY 1 LGCTITSL 7
||| | |
Db 48 LGCTITSL 54
||| | |
RESULT 9
H86372
protein F508.40 [imported] - Arabidopsis thaliana
C:Species: Arabidopsis thaliana (mouse-ear cress)
C>Date: 02-Mar-2001 #sequence_revision 02-Mar-2001 #text_change 31-Mar-2001
C:Accession: H86372
R:Theologis, A.; Ecker, J.R.; Palm, C.J.; Federspiel, N.A.; Kaul, S.; White, C.; A
Chen, C.W.; Chung, M.K.; Conn, L.; Conway, A.B.; Huizar, L.
Nature 408, 816-820, 2000
A:Authors: Hunter, J.L.; Jenkins, J.; Johnson-Hopson, C.; Khan, S.; Khaykin, E.; K
C.A.; Li, J.H.; Li, Y.; Lin, X.; Liu, S.X.; Liu, Z.A.; Luros, J.S.; Maiti, R.; Mar
Rizzo, M.; Rooney, T.; Rowley, D.; Sakano, H.
A:Authors: Salzberg, S.L.; Schwartz, J.R.; Shinn, P.; Southwick, A.M.; Sun, H.; Ta
ker, M.; Wu, D.; Yu, G.; Fraser, C.M.; Venter, J.C.; Davis, R.W.
A:Title: Sequence and analysis of chromosome 1 of the plant Arabidopsis.
A:Reference number: A86141; MUID:21016719; PMID:11130712
A:Accession: H86372
A>Status: preliminary
A:Molecule type: DNA
A:Residues: 1-338 <STC>
A:Cross-references: GB:AA005172; NID:94056465; PIDN:AA096039.1; GSPDB:GN00141
C:Genetics:
A:Gene: F508.40
A:Map position: 1
Query Match 77.3% Score 34; DB 2; Length 338;
Best Local Similarity 85.7% Pred. No. 83;
Matches 6; Conservative 1; Mismatches 0; Indels 0; Gaps 0;
QY 3 GCLITSL 9
||| | | |
Db 136 GCLITSL 142
||| | | |
RESULT 10
S86372
protein F508.39 [imported] - Arabidopsis thaliana
C:Species: Arabidopsis thaliana (mouse-ear cress)
C>Date: 02-Mar-2001 #sequence_revision 02-Mar-2001 #text_change 31-Mar-2001
C:Accession: S86372
R:Theologis, A.; Ecker, J.R.; Palm, C.J.; Federspiel, N.A.; Kaul, S.; White, C.; A
Chen, C.W.; Chung, M.K.; Conn, L.; Conway, A.B.; Huizar, L.
Nature 408, 816-820, 2000
A:Authors: Hunter, J.L.; Jenkins, J.; Johnson-Hopson, C.; Khan, S.; Khaykin, E.; K
C.A.; Li, J.H.; Li, Y.; Lin, X.; Liu, S.X.; Liu, Z.A.; Luros, J.S.; Maiti, R.; Mar
Rizzo, M.; Rooney, T.; Rowley, D.; Sakano, H.
A:Authors: Salzberg, S.L.; Schwartz, J.R.; Shinn, P.; Southwick, A.M.; Sun, H.; Ta
ker, M.; Wu, D.; Yu, G.; Fraser, C.M.; Venter, J.C.; Davis, R.W.
A:Title: Sequence and analysis of chromosome 1 of the plant Arabidopsis.
A:Reference number: A86141; MUID:21016719; PMID:11130712
A:Accession: S86372
A>Status: preliminary
A:Molecule type: DNA
A:Residues: 1-345 <STO>
A:Cross-references: GB:AE005172; NID:94056466; PIDN:AA096039.1; GSPDB:GN00141

C:Genetics:

A:Gene: F508.35

A:Map position: 1

Query Match 77.3%; Score 34; DB 2; Length 345;

Best Local Similarity 85.7%; Pred. No. 85;

Matches 5; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 3 LGCIITS 9

II:II:II

Db 146 LGCIITS 152

RESULT 11

PC4163

toxin-co-regulated protein chain 1 - Vibrio cholerae (fragment)

C:Species: Vibrio cholerae

C:Date: 10-May-1996 #sequence_revision 19-Jul-1996 #text_change 17-Nov-2000

C:Accession: PC4163; S23261

R:Oglerman, M.A.; Voss, E.; Meaney, C.; Faust, R.; Attridge, S.R.; Manning, P.A.

Gene 170, 9-16, 1996

A:Title: Comparison of the promoter proximal regions of the toxin-co-regulated top gene

A:Reference number: JC4719; MUID:96200845; PMID:8621056

A:Accession: PC4163

A:Molecule type: DNA

A:Residues: 1-428 <OGI>

A:Cross-references: EMBL:X64098

R:Manning, P.A.

submitted to the EMBL Data Library, January 1992

A:Reference number: S23261

A>Status: preliminary

A:Molecule type: DNA

A:Residues: 1-401 /RRPKN/ <MAN>

A:Cross-references: EMBL:X64098; NID:q48404; PIDN:CAA45452.1; EID:q48405

C:Comment: This is a methyl-accepting chemotaxis protein.

C:Genetics:

A:Gene: top1

Query Match

Best Local Similarity 87.5%; Score 34; DB 2; Length 428;

Matches 7; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 LGCIITS 8

II:II:II

Db 11 LGCIITS 18

RESULT 12

S48255

probable membrane protein YBR086c - yeast (Saccharomyces cerevisiae)

N:Alternate names: hypothetical protein YBR086c

C:Species: Saccharomyces cerevisiae

C:Date: 03-Aug-1995 #sequence_revision 11-Aug-1995 #text_change 19-Apr-2002

C:Accession: S48255; S45954; S44670

R:Mannhaupt, G.; Stocker, R.; Ehle, S.; Vetter, L.; Feldmann, H.

Yeast 10, 1363-1381, 1994

A:Title: Analysis of a 70 kb region on the right arm of yeast chromosome II.

A:Reference number: S48255; MUID:95208357; PMID:7407426

A:Accession: S48255

A>Status: nucleic acid sequence not shown

A:Molecule type: DNA

A:Residues: 1-946 <MAN>

A:Cross-references: EMBL:X78993; NID:q475045; PIDN:CAA55593.1; PID:q476042

R:Feldmann, H.; Mannhaupt, G.; Schwarze, C.; Vetter, L.

submitted to the Protein Sequence Database, August 1994

A:Reference number: S45927

A:Accession: S45954

A:Molecule type: DNA

A:Residues: 1-946 <FF2>

A:Cross-references: EMBL:X78993; NID:q35351; PID:q536352; MIPS:YBR086c

C:Genetics:

A:Gene: SGD:IST2

A:Cross-references: SGD:S0000290

A:Map position: 2R

C:Superfamily: Saccharomyces cerevisiae probable membrane protein YBR086c

C:Keywords: transmembrane protein

F:131-147/Domain: transmembrane #status predicted <TM1>

F:158-174/Domain: transmembrane #status predicted <TM2>

F:207-243/Domain: transmembrane #status predicted <TM3>

F:248-274/Domain: transmembrane #status predicted <TM4>

F:302-324/Domain: transmembrane #status predicted <TM5>

F:450-477/Domain: transmembrane #status predicted <TM6>

F:506-532/Domain: transmembrane #status predicted <TM7>

F:563-588/Domain: transmembrane #status predicted <TM8>

Query Match

Best Local Similarity 77.3%; Score 34; DB 2; Length 946;

Matches 5; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 LGCIITS 5

II:II:II

Db 520 LGCIITS 527

RESULT 13

T31431

inositol 1,4,5-trisphosphate receptor, localized in plasma membrane - Panulirus argus

C:Species: Panulirus argus

C:Date: 19-May-2000 #sequence_revision 19-May-2000 #text_change 19-May-2000

C:Accession: T31431

R:Munger, S.D.; Ache, B.W.; Greenberg, R.M.

submitted to the EMBL Data Library, March 1998

A:Description: Plasma membrane localization of an olfactory inositol 1,4,5-trisphos

A:Reference number: Z21030

A:Accession: T31431

A>Status: preliminary

A:Molecule type: mRNA

A:Residues: 1-2783 <MUN>

A:Cross-references: EMBL:AF055079; NID:q3660666; PID:q3660667; PIDN:AAC61691.1

C:Superfamily: inositol-trisphosphate receptor

Query Match

Best Local Similarity 77.3%; Score 34; DB 2; Length 2783;

Matches 7; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 LGCIITS 9

II:II:II

Db 2554 ELMCIIVSL 2562

RESULT 14

S00755

platelet/leukocyte C kinase substrate (pieckstrin)

N:Alternate names: p47; platelet/leukocyte C kinase substrate (pieckstrin)

C:Species: Homo sapiens (man)

C:Date: 31-Dec-1988 #sequence_revision 31-Dec-1988 #text_change 20-Apr-2000

C:Accession: S00755; A45762

R:Tyers, M.; Rachubinski, R.A.; Stewart, M.J.; Varrichio, A.M.; Short, R.G.D.; Hasl,

Nature 333, 470-473, 1988

A:Title: Molecular cloning and expression of the major protein kinase C substrate o

A:Reference number: S00755; MUID:88232910; PMID:2897630

A:Accession: S00755

A:Molecule type: mRNA

A:Residues: 1-350 <TYE>

A:Cross-references: EMBL:X07743; NID:q35517; PIDN:CAA30564.1; PID:q35518

R:Tyers, M.; Haslam, R.J.; Rachubinski, R.A.; Hatley, C.B.

J. Cell. Biochem. 40, 133-145, 1989

A:Title: Molecular analysis of pieckstrin: the major protein kinase C substrate of i

A:Reference number: A45762; MUID:89359547; PMID:2768345

A:Accession: A45762

A:Molecule type: mRNA

A:Residues: 1-350 <TYE>

A:Cross-references: GB:X07743; NID:q35517; PIDN:CAA30564.1; PID:q35518

A:Note: 92-Arg was also found

C:Genetics:

A:Gene: GDB:PLEK; p47
A:Cross-references: GDB:383661s; OMIM:174570
A:Map position: 2pter-2qter
A:Superfamily: pleckstrin; pleckstrin repeat homology
C:Keywords: phosphoprotein; signal transduction
F:3-99/Domain: pleckstrin repeat homology <PLK1>
F:243-345/Domain: pleckstrin repeat homology <PLK2>

Query Match 75.04; Score 33; DB 1; Length 350;
Best Local Similarity 55.6%; Pred. No. 1.3e+02;
Matches 5; Conservative 3; Mismatches 1; Indels 0; Gaps 0;

QY 1 LLGCIITSL 9
Db 292 LRGCVVTSV 308
1
1

RESULT 15
D83311
conserved hypothetical protein FA2562 [imported] - Pseudomonas aeruginosa (strain PA01)
C:Species: Pseudomonas aeruginosa
C>Date: 15-Sep-2000 #sequence_revision 15-Sep-2000 #text_change 31-Dec-2000
C:Accession: D83311
R:Stover, C.K.; Pham, X.Q.; Erwin, A.L.; Mizoguchi, S.D.; Warren, P.; Hickey, M.J.; Br
adman, S.; Yuan, Y.; Brody, L.L.; Collier, S.N.; Folger, K.R.; Kas, A.; Larbig, K.; Lim
; Lory, S.; Olson, M.V.
Nature 406, 959-964, 2003
A:Title: Complete genome sequence of Pseudomonas aeruginosa PA01, an opportunistic patho
A:Reference number: A82950; M31D:20437337; PMID:12694012
A:Accession: D83311
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-307 <STO>
A:Cross-references: GH:A82004895; GH:A82004893; NID:q944870; PION:AAG 6650.1; GSPH:HGNC0
A:Experimental source: strain PA01
C:Genetics:
A:Gene: PA2562

Query Match 75.04; Score 33; DB 2; Length 397;
Best Local Similarity 55.6%; Pred. No. 1.5e+02;
Matches 5; Conservative 3; Mismatches 1; Indels 0; Gaps 0;

QY 1 LLGCIITSL 9
Db 221 LLGCVLVAL 229
1
1

Search completed: September 29, 2003, 19:05:56
Job time : 22 secs

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: September 29, 2003, 18:58:47 ; Search time 10 Seconds
(without alignments)
42.324 Million cell updates/sec

Title: US-09-965-594-1_COPY_14_22
Perfect score: 44
Sequence: 1 LSCCHHS 9

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 127863 seqs, 4726735 residues

Total number of hits satisfying chosen parameters: 127859

Minimum DB seq length: 0
Maximum DB seq length: 20000000
Post-processing: Minimum Match 0%
Maximum Match 99%
Listing first 45 summaries

Database : Swissprot_41*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	43	97.7	310	1	POLG_HCVJT
2	40	90.5	310	1	POLG_HCVJT
3	36	81.8	300	1	T289_MOUSE
4	35	79.5	653	1	EGN_PIG
5	34	77.3	946	1	YB76_YEAST
6	33	75.0	318	1	T287_HUMAN
7	33	75.0	350	1	PLEK_HUMAN
8	33	75.0	350	1	PLEK_MOUSE
9	33	75.0	406	1	ARGO_PSEPK
10	33	75.0	437	1	FLHF_PSEPK
11	33	75.0	441	1	YB8E_BACSU
12	33	75.0	795	1	SYFB_SHEON
13	33	75.0	1089	1	POLG_HUMAN
14	33	75.0	1906	1	DICE_MOUSE
15	33	75.0	1912	1	DICE_HUMAN
16	32	72.7	222	1	PSD9_RAT
17	32	72.7	301	1	RECS_TODPA
18	32	72.7	330	1	KUGT_BACSU
19	32	72.7	398	1	YJGN_ECOLI
20	32	72.7	411	1	RAPS_CHICK
21	32	72.7	419	1	AGAL_MOUSE
22	32	72.7	531	1	GHT2_SCHPO
23	32	72.7	689	1	TGM2_CAVCU
24	32	72.7	928	1	YDGI_SCHPO
25	32	72.7	1074	1	SM5A_HUMAN
26	32	72.7	1077	1	SM5A_MOUSE
27	31	70.5	148	1	SSRP_NEIMA
28	31	70.5	162	1	ILJ5_BOVIN
29	31	70.5	180	1	SP22_MOUSE
30	31	70.5	237	1	LAPT_PASHA
31	31	70.5	312	1	MCAL_HUMAN
32	31	70.5	312	1	T285_MOUSE
33	31	70.5	312	1	YM17_YEAST

34	31	70.5	334	1	YC52_PSEAE
35	31	70.5	398	1	YJGN_ECOLI
36	31	70.5	411	1	RAPS_HUMAN
37	31	70.5	411	1	RAPS_MOUSE
38	31	70.5	411	1	RAPS_TORCA
39	31	70.5	417	1	PCL_RHOCA
40	31	70.5	427	1	ETLR_PIG
41	31	70.5	436	1	Y326_METJA
42	31	70.5	439	1	IOLF_BACSU
43	31	70.5	495	1	POLG_DEN23
44	31	70.5	556	1	PTI_STRCO
45	31	70.5	609	1	THI3_YEAST

ALIGNMENTS

RESULT 1
POLG_HCVJT STANDARD: PRT: 301C AA.
AC QGQ263;
BT 01-APR-1993 (Rel. 25, Last sequence update)
DT 01-APR-1993 (Rel. 25, Last annotation update)
DE Genome polypeptide [Contains: Capsid protein C (Core protein) (P22); Envelope glycoprotein E1 (GP32) (GP35); Envelope glycoprotein E2 (GP68) (GP70) (NS1); Protein P7; Nonstructural protein NS2 (P21) (EC 3.4.22.-); Protease/helicase NS3 (P70) (Hepacivirus) (EC 3.4.21.98); Nonstructural protein NS4A (P4); Nonstructural protein NS4B (P66) (P70) (RNA-directed RNA polymerase) (EC 2.7.7.48)].
DE Hepatitis C virus (isolate HC-JT) (HCV).
OC Viruses; ssRNA positive-strand viruses, no DNA stage: Flaviviridae;
OC Hepacivirus.
OX NCBI_TaxID=31642;
RN 111
RF SEQUENCE FROM N.A.
RX MEDLINE=92295714; PubMed=1318527;
SA Tanaka T., Kato N., Nakagawa M., Ootsuyama Y., Cho M.J., Nakazawa T., Hilkata M., Ishimura Y., Shimotohno K.;
RT "Molecular cloning of hepatitis C virus genome from a single Japanese carrier: sequence variation within the same individual and among infected individuals."
RI Virus Res. 23:39-53(1992)
CC 1- FUNCTION: THE SMALL PROTEINS NS2A, NS2B, NS4A AND NS4B ARE HYDROPHOBIC, SUGGESTING A POSSIBLE MEMBRANE-RELATED FUNCTION. NS3 AND NS5 MAY PLAY A ROLE IN THE VIRAL RNA REPLICATION.
CC 2- CATALYTIC ACTIVITY: Hydrolysis of four peptide bonds in the viral precursor polypeptide, commonly with Asp or Glu in the P6 position, Cys or Thr in P1 and Ser or Ala in P1'.
CC 3- CATALYTIC ACTIVITY: N nucleoside triphosphate = N diphosphate + {RNA}(N).
CC 4- SUBUNIT: THE VIRION OF THIS VIRUS IS A NUCLEOCAPSID COVERED BY A LIPOPROTEIN ENVELOPE. THE ENVELOPE CONSISTS OF TWO PROTEINS: PROTEIN M AND RNA.
CC 5- SIMILARITY: THE PROTEASE BELONGS TO PEPTIDASE FAMILY S29.
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CC
CC EMBL: D11168; BAA01943.1;
CC PIR: A45573; A45573
CC PDB: 1A1Q; 25-MAR-96.
CC PDB: 1JXP; 14-JAN-98.
CC MEROPS: S29.001;
CC MEROPS: U39.001;
CC InterPro: IPR001410; DEAD.

DR InterPro: IPR002522; HCV_capsid.
 DR InterPro: IPR002521; HCV_core.
 DR InterPro: IPR002519; HCV_env.
 DR InterPro: IPR002511; HCV_NS1.
 DR InterPro: IPR002518; HCV_NS2.
 DR InterPro: IPR004109; HCV_NS3.
 DR InterPro: IPR00745; HCV_NS4a.
 DR InterPro: IPR002493; HCV_NS4b.
 DR InterPro: IPR002869; HCV_NS5a.
 DR InterPro: IPR002166; HCV_RdRp.
 DR InterPro: IPR007095; RNA_pol_DS_PS.
 DR InterPro: IPR007094; RNA_pol_PSVir.
 DR Pfam: PF01543; HCV_core; 1.
 DR Pfam: PF01542; HCV_core; 1.
 DR Pfam: PF01539; HCV_env; 1.
 DR Pfam: PF01560; HCV_NS1; 1.
 DR Pfam: PF01538; HCV_NS2; 1.
 DR Pfam: PF02907; HCV_NS3; 1.
 DR Pfam: PF01006; HCV_NS4a; 1.
 DR Pfam: PF01001; HCV_NS4b; 1.
 DR Pfam: PF01506; HCV_NS5a; 1.
 DR Pfam: PF00271; Helicase; 1.
 DR Pfam: PF00998; Viral_RdRp; 1.
 DR Pfam: PF086962; HCV_NS1; 1.
 DR Pfam: SMO487; DEXdc; 1.
 DR PolyProtein: Glycoprotein; Transferrase; RNA-directed RNA polymerase;
 KW Core protein; Coat protein; Envelope protein; Helicase; AIP-binding;
 KW Transmembrane; Nonstructural protein; Hydrolase; Serine protease;
 KW 3D-structure.
 FT INIT_MET 1
 FT CHAIN 1 125
 FT CHA-N 116 191
 FT CHA-N 192 383
 FT CHA-N 384 729
 FT CHA-N 730 1006
 FT CHA-N 1007 1615
 FT CHA-N 1616 1862
 FT CHA-N 1863 2013
 FT CHA-N 2014 3010
 FT CHA-N 3010 369
 FT TRANSMEM 347 369
 FT ACT_SITE 1083 1083
 FT ACT_SITE 1107 1107
 FT ACT_SITE 1165 1165
 FT NP_BIND 1240 1237
 FT SITE 1316 1319
 FT CARBOHYD 196 196
 FT CARBOHYD 209 209
 FT CARBOHYD 234 234
 FT CARBOHYD 250 250
 FT CARBOHYD 305 305
 FT CARBOHYD 417 417
 FT CARBOHYD 423 423
 FT CARBOHYD 430 430
 FT CARBOHYD 448 448
 FT CARBOHYD 532 532
 FT CARBOHYD 540 540
 FT CARBOHYD 556 556
 FT CARBOHYD 576 576
 FT CARBOHYD 623 623
 FT CARBOHYD 645 645
 FT CARBOHYD 2041 2041
 FT CARBOHYD 2077 2077
 FT CARBOHYD 2246 2246
 FT CARBOHYD 2523 2523
 FT CARBOHYD 2788 2788
 FT CARBOHYD 3010 AA; 326573 MW; 94A1C77435D642B5 CRC64;
 Query Match: 97.7%; Score 43; DR 1; Length 3010;
 Best Local Similarity 86.9%; Pred. No. 5.9;
 Matches 8; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 LLGCIITSL 9

DB 1039 LLGCIITSL 1047
 RESULT 2
 ID POLA_HCVTW STANDARD: PRT: 3010 AA.
 AC P29846;
 DT 01-APR-1993 (Rel. 25, Created)
 DT 01-APR-1993 (Rel. 25, Last annotation update)
 DT 15-SEP-2003 (Rel. 42, Last annotation update)
 DE Genome polyprotein. Contains: Capsid protein C (Core protein) (P22);
 DE Envelope glycoprotein E1 (GP32) (GP35); Envelope glycoprotein E2
 DE (GP68) (GP70) (NS1); Protein P7; Nonstructural protein NS2 (P21)
 DE (EC 3.4.22.-); Protease/helicase NS3 (P70) (Hepacivirin)
 DE (EC 3.4.21.98); Nonstructural protein NS4A (P4); Nonstructural protein
 DE NS4B (P27); Nonstructural protein NS5A (P56); Nonstructural protein
 DE NS5B (P66) (P70) (RNA-directed RNA polymerase) (EC 2.7.7.48)].
 OS Hepatitis C virus (isolate Taiwan) (HCV).
 OC Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
 OC Hepacivirus.
 KX NCBI_TaxID-31645;
 RN 1
 RP SEQUENCE FROM N.A.
 RX MEDLINE-92230206; PubMed-1314449;
 KA Chen P.J., Lin M.H., Tai K.F., Liu P.C., Lin C.J., Chen D.S.;
 FT "The Taiwanese hepatitis C virus genome: sequence determination and
 FT mapping the 5' termini of viral genomic and antigenomic RNA.";
 FT Virology 188:102-113(1992).
 CC -!- FUNCTION: THE SMALL PROTEINS NS2A, NS2B, NS4A AND NS4B ARE
 CC HYDROPHOBIC, SUGGESTING A POSSIBLE MEMBRANE-RELATED FUNCTION.
 CC NS3 AND NS5 MAY PLAY A ROLE IN THE VIRAL RNA REPLICATION.
 CC -!- CATALYTIC ACTIVITY: Hydrolysis of four peptide bonds in the viral
 CC precursor polyprotein, commonly with asp or glu in the P6
 CC position, Cys or Thr in P1 and Ser or Ala in P1'.
 CC -!- CATALYTIC ACTIVITY: N nucleoside triphosphate - N diphosphate +
 CC [RNA](N).
 CC -!- SUBUNIT: THE VIRION OF THIS VIRUS IS A NUCLEOCAPSID COVERED BY A
 CC LIPOPROTEIN ENVELOPE. THE ENVELOPE CONSISTS OF TWO PROTEINS:
 CC PROTEIN M AND GPROTEIN E. THE NUCLEOCAPSID IS A COMPLEX OF
 CC PROTEIN C AND MRNA.
 CC -!- SIMILARITY: THE PROTEASE BELONGS TO PEPTIDASE FAMILY S29.
 CC
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 CC
 CC EMBL: M84754; ; NOT_ANNOTATED_CDS.
 CC PIR: A40244; GNAVTV
 CC PDB: 1N64; 25-FEB-03.
 CC PDB: 1NS3; 08-APR-98.
 CC MEROPS: S29.001; -.
 CC MESOPS: G39.001; -.
 CC InterPro: IPR001410; DEAD.
 CC InterPro: IPR002522; HCV_capsid.
 CC InterPro: IPR002521; HCV_core.
 CC InterPro: IPR002519; HCV_env.
 CC InterPro: IPR00253; HCV_NS1.
 CC InterPro: IPR002518; HCV_NS2.
 CC InterPro: IPR004109; HCV_NS3.
 CC InterPro: IPR00745; HCV_NS4a.
 CC InterPro: IPR001490; HCV_NS4b.
 CC InterPro: IPR002868; HCV_NS5a.
 CC InterPro: IPR002166; HCV_RdRp.
 CC InterPro: IPR007095; RNA_pol_DS_PS.
 CC InterPro: IPR007094; RNA_pol_PSVir.
 CC Pfam: PF01543; HCV_core; 1.
 CC Pfam: PF01542; HCV_core; 1.
 CC Pfam: PF01539; HCV_env; 1.

DR	Pfam: PF01560; HCV_NSI; 1.	US	Mus musculus (Mouse).
DR	Pfam: PF01538; HCV_NS2; 1.	OC	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
DR	Pfam: PF02907; HCV_NS3; 1.	OC	Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
DR	Pfam: PF01006; HCV_NS4a; 1.	OX	NCBI_taxonomy:10090;
DR	Pfam: PF01001; HCV_NS4b; 1.	RN	[1]
DR	Pfam: PF01506; HCV_NS5a; 1.	RP	SEQUENCE FROM N.A., AND TOPOLOGY.
DR	Pfam: PF00271; Helicase_C; 1.	RC	STRAIN-129/Svj;
DR	Pfam: PF00998; Viral_RdRP; 1.	RX	MEDLINE-20222571; PubMed-10761934;
DR	ProDom: PD186062; HCV_NS1; 1.	RA	Adler E., Hoon M.A., Mueller K.L., Chandrashekar J., Ryba N.J.P.,
DR	SMART: SM00487; DEXDC; 1.	RA	Zuker C.S.;
KW	Polyprotein; Glycoprotein; Transferase; RNA-directed RNA polymerase;	RI	"A novel family of mammalian taste receptors.";
KW	Core protein; Coat protein; Envelope protein; Helicase; ATP-binding;	RL	Cell 100:693-702(2000).
KW	Transmembrane; Nonstructural protein; Hydrolase; Serine protease;	RN	[2]
KW	3D-structure.	RP	CHARACTERIZATION.
FI	INIT_MEI 1	RX	MEDLINE-20222572; PubMed-10761935;
FI	CHAIN 1	RA	Chandrashekar J., Mueller K.L., Hoon M.A., Adler E., Peng L., Guo W.,
FI	CHAIN 116	RA	Zuker C.S., Ryba N.J.;
FI	CHAIN 191	RI	"T2Rs function as bitter taste receptors.";
FI	CHAIN 192	RL	Cell 100:703-711(2000).
FI	CHAIN 384	RN	[3]
FI	CHAIN 729	RP	REVIEW.
FI	CHAIN 730	RX	MEDLINE-22135574; PubMed-12139982;
FI	CHAIN 1007	RA	Montmayeur J.-P., Matsunami H.;
FI	CHAIN 1615	RI	"Receptors for bitter and sweet taste.";
FI	CHAIN 1862	RL	Curr. Opin. Neurobiol. 12:366-371(2002).
FI	CHAIN 1863	RN	[4]
FI	CHAIN 2014	RP	REVIEW.
FI	CHAIN 2014	RX	MEDLINE-21634924; PubMed-11696554;
FI	CHAIN 347	RA	Margolskee R.F.;
FI	CHAIN 362	RI	"Molecular mechanisms of bitter and sweet taste transduction.";
FI	ACT_SITE 1063	RL	J. Biol. Chem. 277:1-4(2002).
FI	ACT_SITE 1107	RN	[5]
FI	ACT_SITE 1165	RP	REVIEW.
FI	ACT_SITE 1230	RX	MEDLINE-22469025; PubMed-12581520;
FI	NP_BIND 1230	RA	Zhang Y., Hoon M.A., Chandrashekar J., Mueller K.L., Cook B., Wu D.,
FI	SITE 1315	RI	Zuker C.S., Ryba N.J.;
FI	N-LINKED (GLCNAC) 196	RL	"Coding of sweet, bitter, and umami tastes: different receptor cells
FI	N-LINKED (GLCNAC) 209	RN	sharing similar signaling pathways.";
FI	N-LINKED (GLCNAC) 233	RP	Cell 112:293-301(2003).
FI	N-LINKED (GLCNAC) 234	CC	FUNCTION: Receptor that may play a role in the perception of
FI	N-LINKED (GLCNAC) 250	CC	bitterness and is gustducin-linked. May play a role in sensing the
FI	N-LINKED (GLCNAC) 305	CC	chemical composition of the gastrointestinal content. The activity
FI	N-LINKED (GLCNAC) 305	CC	of this receptor may stimulate alpha gustducin, mediate PLC-beta-2
FI	N-LINKED (GLCNAC) 417	CC	activation and lead to the gating of TRPM5. Functions as a ligand
FI	N-LINKED (GLCNAC) 423	CC	for cyclotriamide.
FI	N-LINKED (GLCNAC) 423	CC	SUBCELLULAR LOCATION: Integral membrane protein.
FI	N-LINKED (GLCNAC) 430	CC	TISSUE SPECIFICITY: Expressed in subsets of taste receptor cells
FI	N-LINKED (GLCNAC) 448	CC	of the tongue and palate epithelium and exclusively in gustducin-
FI	N-LINKED (GLCNAC) 532	CC	positive cells. Expressed in gastric and duodenal tissues.
FI	N-LINKED (GLCNAC) 540	CC	MISCELLANEOUS: Most taste cells may be activated by a limited
FI	N-LINKED (GLCNAC) 556	CC	number of bitter compounds; individual taste cells can
FI	N-LINKED (GLCNAC) 576	CC	discriminate among bitter stimuli.
FI	N-LINKED (GLCNAC) 623	CC	SIMILARITY: Belongs to family T2R of G-protein coupled receptors.
FI	N-LINKED (GLCNAC) 645	CC	CAUTION: This protein was previously referred to T2R5, but due to
FI	N-LINKED (GLCNAC) 2041	CC	its high similarity towards both the human and rat T2R9 sequences
FI	N-LINKED (GLCNAC) 2077	CC	it is considered to be T2R9.
FI	N-LINKED (GLCNAC) 2240	CC	-----
FI	N-LINKED (GLCNAC) 2529	CC	This SWISS-PROT entry is copyright. It is produced through a collaboration
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FI	N-LINKED (GLCNAC) 2788	CC	the European Bioinformatics Institute. There are no restrictions on its
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FI	N-LINKED (GLCNAC) 3010	CC	entities requires a license agreement (See http://www.isb-sib.ch/announce/
FI	N-LINKED (GLCNAC) 3010	CC	or send an email to license@isb-sib.ch).
FI	N-LINKED (GLCNAC) 3010	CC	-----
FI	N-LINKED (GLCNAC) 3010	CC	EMBL: AF227147; AAF43920.1;
FI	N-LINKED (GLCNAC) 3010	CC	Pfam: PF05296; TAS2R; 1.
FI	N-LINKED (GLCNAC) 3010	CC	Receptor: G-protein coupled receptor; Transmembrane.
FI	N-LINKED (GLCNAC) 3010	CC	DOMAIN 1 7 EXTRACELLULAR (POTENTIAL).
FI	N-LINKED (GLCNAC) 3010	CC	TRANSMEM 8 28 1 (POTENTIAL).
FI	N-LINKED (GLCNAC) 3010	CC	DOMAIN 29 43 CYTOPLASMIC (POTENTIAL).
FI	N-LINKED (GLCNAC) 3010	CC	TRANSMEM 44 64 2 (POTENTIAL).
FI	N-LINKED (GLCNAC) 3010	CC	DOMAIN 65 87 EXTRACELLULAR (POTENTIAL).
QY	1 LLGCITSL 9	CC	-----
DB	1039 LFGCITSL 1047	CC	-----
RESULT 3			
T2R9_MOUSE			
ID	T2R9_MOUSE	STANDARD:	PRT: 300 AA.
AC	Q9JKT4;		
DT	15-SEP-2003 (Rel. 42, Created)		
DT	15-SEP-2003 (Rel. 42, Last sequence update)		
DT	15-SEP-2003 (Rel. 42, Last annotation update)		
DE	Taste receptor type 2 member 9 (T2R9) (Taste receptor type 2 member		
DE	5) (T2R5).		
GN	TAS2R5.		

Query Match 90.9%; Score 40; DB 1; Length 3010;
 Best Local Similarity 88.9%; Pred. No. 21;
 Matches 8; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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FT TRANSMEM 88 108 3 (POTENTIAL)
FT DOMAIN 109 128 CYTOPLASMIC (POTENTIAL)
FT TRANSMEM 129 149 4 (POTENTIAL)
FT DOMAIN 150 181 EXTRACELLULAR (POTENTIAL)
FT TRANSMEM 182 202 5 (POTENTIAL)
FT DOMAIN 203 233 CYTOPLASMIC (POTENTIAL)
FT TRANSMEM 234 254 6 (POTENTIAL)
FT DOMAIN 255 259 EXTRACELLULAR (POTENTIAL)
FT TRANSMEM 260 280 7 (POTENTIAL)
FT DOMAIN 281 303 CYTOPLASMIC (POTENTIAL)
FT CARBOHYD 151 161 N-LINKED (GLCNAC...) (POTENTIAL)
SQ SEQUENCE 300 AA: 34416 MW: 92AD449C66FB85EB CRC64;

Query Match 81.8%; Score 36; D3 1; Length 300;
Best Local Similarity 75.0%; Pred. No. 15; Mismatches 0; Indels 0; Gaps 0;
Matches 6; Conservative 2;

QY 1 LLGCIITS 8
DB 133 LLGCLTS 140

RESULT 4
EGLN_PIG STANDARD; PRT: 653 AA.
AC P37176;
DT 01-OCT-1994 (Rel. 30, Created)
DI 01-OCT-1994 (Rel. 30, Last sequence update)
DI 28-FEB-2003 (Rel. 41, Last annotation update)
DE Endoglin precursor.
GN ENG.
OS Sus scrofa (Pig).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Cetartiodactyla; Suidae; Suidae; Suidae.
OX NCBI_TaxID=9823;
RN (1)
RP SEQUENCE FROM N.A.
RC TISSUE-Uterus;
RX MEDLINE=94124550; PubMed=8294451;
RA Yamashita H., Ichijo H., Grimsby S., Moren A., ten Dijke P.,
RA Miyazono K.;
RT "Endoglin forms a heteromeric complex with the signaling receptors
RT for transforming growth factor-beta.";
RL J. Biol. Chem. 269:1995-2001(1994).
CC -!- FUNCTION: MAJOR GLYCOPROTEIN OF VASCULAR ENDOTHELIUM. MAY PLAY A
CC CRITICAL ROLE IN THE BINDING OF ENDOTHELIAL CELLS TO INTEGRINS
CC AND/OR OTHER RGD RECEPTORS.
CC -!- SUBUNIT: HOMODIMER THAT FORMS AN HETEROMERIC COMPLEX WITH THE
CC SIGNALING RECEPTORS FOR TRANSFORMING GROWTH FACTOR-BETA: TGF-BETA
CC RECEPTORS 1 AND/OR II. IT IS ABLE TO BIND TGF-BETA 1, AND 3
CC EFFICIENTLY AND TGF-BETA 2 LESS EFFICIENTLY.
CC -!- SUBCELLULAR LOCATION: Type I membrane protein.
CC -!- SIMILARITY: SOME REGIONS OF SIMILARITY TO TGF-BETA RECEPTOR TYPE
CC III (BETAGLYCAN).
CC
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CC
CC EMBL: Z23142; CAA00673.1;
CC PIR: A49722; A49722.
CC InterPro: IPR001507; Endoglin/CD105.
CC Pfam: PF00100; zora_pellucida; 1.
CC SMART: SM00241; 2P; 1.
CC Cell adhesion; Glycoprotein; Transmembrane; Signal.
CC SIGNAL 1 24 POTENTIAL.
CC CHAIN 25 653 ENDOLIN.
CC DOMAIN 25 581 EXTRACELLULAR (POTENTIAL).
CC TRANSMEM 582 606 POTENTIAL.

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FT DOMAIN 607 653 CYTOPLASMIC (POTENTIAL)
FT DOMAIN 335 574 SER/THR-RICH.
FT SITE 396 400 CELL ATTACHMENT SITE (BY SIMILARITY).
FT CARBOHYD 57 57 N-LINKED (GLCNAC...) (POTENTIAL)
FT CARBOHYD 306 306 N-LINKED (GLCNAC...) (POTENTIAL)
SQ SEQUENCE 653 AA: 70279 MW: 7887B4A61D5E3E5A CRC64;

Query Match 79.5%; Score 35; D8 1; Length 653;
Best Local Similarity 87.5%; Pred. No. 46; Mismatches 0; Indels 0; Gaps 0;
Matches 7; Conservative 0;

QY 2 LGCIITSL 9
DB 379 LGCIITSL 386

RESULT 5
YBT6_YEAST STANDARD; PRT: 946 AA.
AC P36250;
DT 01-OCT-1994 (Rel. 30, Created)
DT 01-OCT-1994 (Rel. 30, Last sequence update)
DT 01-NOV-1995 (Rel. 32, Last annotation update)
DE Hypothetical 105.9 kDa protein in AAC3-RFC5 intergenic region.
GN YBC86C OR YBR0809.
OS Saccharomyces cerevisiae (Baker's yeast).
OC Eukaryota; Fungi; Ascomycota; Saccharomycotina; Saccharomycetes;
OC Saccharomycetales; Saccharomycetaceae; Saccharomycos.
OX NCBI_TaxID=4932;
RN (1)
RP SEQUENCE FROM N.A.
RC STRAIN=S288C;
RX MEDLINE=95208357; PubMed=7900426;
RA Mannhaupt G., Stucka R., Ehmele S., Vetter I., Feldmann H.;
RT "Analysis of a 70 kb region on the right arm of yeast chromosome II.";
RL Yeast 10:1363-1381(1994).
CC -!- SUBCELLULAR LOCATION: Integral membrane protein (Potential).
CC
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CC
CC EMBL: X78993; CAA55593.1;
CC EMBL: Z35955; CAA85034.1;
CC PIR: S48255; S48255.
CC SGD: S0000290; IST2.
CC Pfam: PF04547; DUF590; 1.
CC Hypothetical protein; Transmembrane.
CC TRANSMEM 122 142 POTENTIAL.
CC TRANSMEM 154 174 POTENTIAL.
CC TRANSMEM 218 238 POTENTIAL.
CC TRANSMEM 254 274 POTENTIAL.
CC TRANSMEM 303 323 POTENTIAL.
CC TRANSMEM 448 468 POTENTIAL.
CC TRANSMEM 506 526 POTENTIAL.
CC TRANSMEM 564 584 POTENTIAL.
CC SEQUENCE 946 AA: 105903 MW: F51A43A5D378B7BC CRC64;

Query Match 77.3%; Score 34; D5 1; Length 946;
Best Local Similarity 62.5%; Pred. No. 96; Mismatches 5; Conservative 3;

QY 1 LLGCIITS 8
DB 520 LLGCVITA 527

RESULT 6
T2R7_HUMAN

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ID T2R7_HUMAN STANDARD; PRI: 318 AA.
 AC Q9NYW3;
 DT 15-SEP-2003 (Rel. 42, Last sequence update);
 DT 15-SEP-2003 (Rel. 42, Last sequence update);
 DE Taste receptor type 2 member 7 (T2R7) (Taste receptor family B member 4) (T2R4).
 DE TAS2R7.
 GN Homo sapiens (Human).
 OS Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 OX NCBI_TaxID=9606;
 RN [1]
 RP SEQUENCE FROM N.A., AND TECHNOLOGY.
 RX MEDLINE=20222571; PubMed=10761934;
 RA Adler E., Hoon M.A., Mueller K.L., Chandrasekhar J., Ryba N.J.P., Zuker C.S.;
 RA "A novel family of mammalian taste receptors."; Cell 100:693-702(2000).
 RL [2]
 RP CHARACTERIZATION.
 RX MEDLINE=20222572; PubMed=10761935;
 RA Chandrasekhar J., Mueller K.L., Hoon M.A., Adler E., Feng L., Guo W., Zuker C.S., Ryba N.J.;
 RA "T2Rs function as bitter taste receptors."; Cell 100:703-711(2000).
 RL [3]
 RP REVIEW.
 RX MEDLINE=22135574; PubMed=12139682;
 RA Montmayeur J.-P., Matsunami H.;
 RA "Receptors for bitter and sweet taste."; Curr. Opin. Neurobiol. 12:366-371(2002).
 RL [4]
 RP REVIEW.
 RX MEDLINE=21634924; PubMed=11566554;
 RA Margolskee R.F.;
 RA "Molecular mechanisms of bitter and sweet taste transduction."; J. Biol. Chem. 277:1-4(2002).
 RL [5]
 RP REVIEW.
 RX MEDLINE=22459025; PubMed=12581520;
 RA Zhang Y., Hoon M.A., Chandrasekhar J., Mueller K.L., Cook R., Wu D., Zuker C.S., Ryba N.J.;
 RA "Coding of sweet, bitter, and umami tastes: different receptor cells sharing similar signalling pathways."; Cell 112:293-301(2003).
 RL [6]
 CC -1- FUNCTION: Receptor that may play a role in the perception of bitterness and is gustducin-linked. May play a role in sensing the chemical composition of the gastrointestinal content. The activity of this receptor may stimulate alpha gustducin, mediate PLC-beta-2 activation and lead to the gating of TRPM5.
 CC -1- SUBCELLULAR LOCATION: Integral membrane protein.
 CC -1- TISSUE SPECIFICITY: Expressed in subsets of taste receptor cells of the tongue and palate epithelium and exclusively in gustducin positive cells.
 CC -1- MISCELLANEOUS: Most taste cells may be activated by a limited number of bitter compounds; individual taste cells can discriminate among bitter stimuli.
 CC -1- SIMILARITY: Belongs to family T2R of G-protein coupled receptors.
 CC
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 CC
 CC EMBL: AF227133; AAF43906.1;
 CC Genbank: HGNC:14913; TAS2R7.
 CC MIM: 604793;
 CC InterPro: IPR000276; GPCR_Rhodpsn.
 CC Pfam: PF05296; TAS2R; 1.

DR PROSITE: PS02622; G-PROTEIN COUPLED RECEPTOR; Transmembrane.
 KW Receptor; G-protein coupled receptor; Transmembrane.
 FT DOMAIN 1 9
 FT TRANSMEM 10 30
 FT DOMAIN 31 55
 FT TRANSMEM 56 76
 FT DOMAIN 77 94
 FT TRANSMEM 95 115
 FT DOMAIN 116 128
 FT TRANSMEM 129 149
 FT DOMAIN 150 187
 FT TRANSMEM 188 208
 FT DOMAIN 209 235
 FT TRANSMEM 236 256
 FT DOMAIN 257 266
 FT TRANSMEM 267 287
 FT DOMAIN 288 318
 FT CARBOHYD 167 167
 FT CARBOHYD 175 175
 SQ SEQUENCE 318 AA: 36549 MW: F187ADB2D8274B8A CRC64;
 Query Match 75.0%; Score 33; DB 1; Length 318;
 Best Local Similarity 55.6%; Pred. No. 59;
 Matches 5; Conservative 3; Mismatches 1; Indels 0; Gaps 0;
 QY 1 LLGCIITSL 9
 DI 134 LLGCVVLSV 142
 AS-SU1 7
 PLEK_HUMAN STANDARD; PRI: 350 AA.
 AC P05567; Q8WV61;
 DI C1-AUG-1988 (Rel. 38, Created)
 DI C1-AUG-1988 (Rel. 38, Last sequence update)
 DT 15-SEP-2003 (Rel. 42, Last annotation update)
 DE Plectstrin (Platelet p47 protein).
 GN PLEK OR P47.
 OS Homo sapiens (Human).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 OX NCBI_TaxID=9606;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=86232920; PubMed=2897530;
 RA Myers M., Rachubinski R.A., McCaw M.L., Varrichio A.M., Shorr R.G.L., Haslam R.J., Harley C.B.;
 RA "Molecular cloning and expression of the major protein kinase C substrate of platelets."; Nature 333:470-473(1988).
 RL [2]
 RN [2]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=89359547; PubMed=2768345;
 RA Myers M., Haslam R.J., Rachubinski R.A., Harley C.B.;
 RA "Molecular analysis of plectstrin: the major protein kinase C substrate of platelets."; J. Cell. Biochem. 40:133-145(1989).
 RL [3]
 RN [3]
 RP SEQUENCE FROM N.A.
 RC TISSUE=Lymph;
 RX MEDLINE=22388257; PubMed=12477932;
 RA Strausberg R.L., Feingold E.A., Grouse L.H., Derge J.G., Klausner R.D., Collins F.S., Wagner L., Shenmen C.M., Schuler G.D., Altschul S.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K., Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Haie F., Diatchenko L., Marusik K., Farmer A.A., Rubin G.M., Hong L., Stapleton M., Soares M.B., Bonaldo M.F., Casavant T.L., Scheetz T.E., Brownstein M.J., Udwin T.B., Toshiyuki S., Carninci P., Prange C., Raha S.S., Loquellano N.A., Peters G.J., Abramson R.D., Mullan S.J., Bosak S.A., McEwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H., Richards S., Worley K.C., Hale S., Garcia A.M., Gay I.J., Hulyk S.W., Villalón D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,

RA Fahey J., Helton E., Kerteman M., Madan A., Rodrigues S., Sanchez A.,
 RA Whiting M., Madan A., Young A.C., Shevchenko Y., Zoufiard G.G.,
 RA Blackley R.W., Grisham J.W., Green E.D., Dickson M.C.,
 RA Rodriguez A.C., Grimwood J., Schmutz J., Myers R.N.,
 RA Butterfield Y.S.N., Krzyzinski M.I., Skalska U., Smallus D.E.,
 RA Scherch A., Schein J.E., Jones S.J.M., Marra M.A.,
 RT *Generation and initial analysis of more than 15,000 full-length
 human and mouse cDNA sequences.*;
 RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).
 RN [4]
 RN STRUCTURE BY NMR OF 1-105.
 RP MEDLINE-9426557; PubMed-8208296;
 RX Yoon H.S., Hajduk P.J., Petros A.M., Olejniczak E.T., Meadows R.P.,
 RA Fesik S.W.,
 RT *Solution structure of a pleckstrin-homology domain.*;
 RL Nature 369:672-675(1994).
 CC EXACT FUNCTION: MAJOR PROTEIN KINASE C SUBSTRATE OF PLATELETS. ITS
 CC -!- FUNCTION: Contains 1 DEP domain.
 CC -!- SIMILARITY: Contains 2 PH domains.
 CC -!- SIMILARITY: Contains 2 PH domains.
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 CC -----
 DR EMBL: X07743; CAA00564.1; .
 DR EMBL: H0018549; AAI18549.1; .
 DR PIR: S00755; SC0755.
 DR PDB: 1PLS; 03-JUN-95.
 DR Genew: HGNC:9070; PLEK.
 DR MIM: 173570; .
 DR GO: GO:0005509; Focal-calcium ion binding activity; TAS.
 DR InterPro: IPR000591; DEP.
 DR Pfam: PF00610; IPR001849; PH.
 DR Pfam: PF00169; PH; 2.
 DR SMART: SM00049; DEP; 1.
 DR SMART: SM00233; PH; 2.
 DR PROSITE: PSS0186; DEP; 1.
 DR PROSITE: PSS0003; PH; DOMAIN; 2.
 KW Phosphorylation; Repeat; 3D-structure.
 FT DOMAIN 4 101
 FT PH 1
 FT DOMAIN 136 221
 FT DEP 2
 FT VARIAT 244 347
 FT W-2, R.
 FT N -> K (IN REF. 3).
 FT CONFLICT 97 97
 FT STRAND 7 14
 FT STRAND 22 29
 FT TURN 30 31
 FT STRAND 32 36
 FT TURN 39 40
 FT TURN 40 42
 FT STRAND 46 49
 FT STRAND 55 56
 FT STRAND 58 73
 FT TURN 74 76
 FT STRAND 77 82
 FT HELIX 86 105
 FT TURN 104 105
 SQ SEQUENCE 350 AA; 40082 MW; 2E2A128CB526361 CRC64;
 Query Match 75.0%; Score 33; DB 1; Length 350;
 Best Local Similarity 55.6%; Pred. No. 63;
 Matches 5; Conservative 3; Mismatches 1; Indels 0; Gaps 0;
 QY 1 LLGC:ITSL 9
 DB 292 LRCCVTVSV 300

RESULT 8
 PLEK_MOUSE STANDARD; PRI: 350 AA.
 AC 09JHK5; 09ERI9;
 DT 28-FEB-2003 (Rel. 41, Created)
 DT 28-FEB-2003 (Rel. 41, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE Pleckstrin.
 GN PLEK.
 OS Mus musculus (Mouse).
 CC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 CC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 CX NCBI_TaxID=10090;
 RN [1]
 RN SEQUENCE FROM N.A.
 RC STRAIN=BALE/C;
 RX MEDLINE-2031822; PubMed-10860665;
 RA Omarik J.L., Hegamyer G., Gerrard B., Dean M., Colburn N.H.,
 RT "cDNA cloning and mapping of mouse pleckstrin (Plek), a gene
 RI upregulated in transformation-resistant cells.*;
 RL Genomics 66:204-212(2000).
 RN [2]
 RN SEQUENCE FROM N.A.
 RA Zhang Y., Wu G., Paige C.J.,
 RT "Involvement of pleckstrin in B cell differentiation and
 RT activation.*;
 RL Submitted (JUN-1998) to the EMBL/GenBank/DBJ databases.
 RN [3]
 RN SEQUENCE FROM N.A.
 RA Ahn H.-J., Cho J.-J.,
 RT "Mouse pleckstrin 1 is induced in mast cells after IgE cross-
 RT linking.*;
 RL Submitted (SEP-2000) to the EMBL/GenBank/DBJ databases.
 CC -!- FUNCTION: MAJOR PROTEIN KINASE C SUBSTRATE OF PLATELETS, ITS
 CC EXACT FUNCTION IS NOT KNOWN.
 CC -!- SIMILARITY: Contains 1 DEP domain.
 CC -!- SIMILARITY: Contains 2 PH domains.
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 CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
 CC or send an email to license@isb-sib.ch).
 CC -----
 DR EMBL: AF181829; AAF75830.1; .
 DR EMBL: AF073294; AAF72039.1; .
 DR EMBL: AF303745; AAG29513.1; .
 DR HSSP: P0H567; IPI5.
 DR MGD: MGI:1860485; Plek.
 DR InterPro: IPR000591; DEP.
 DR InterPro: IPR001849; PH.
 DR Pfam: PF00610; DEP; 1.
 DR Pfam: PF00169; PH; 2.
 DR SMART: SM00049; DEP; 1.
 DR SMART: SM00233; PH; 2.
 DR PROSITE: PSS0186; DEP; 1.
 DR PROSITE: PSS0003; PH; DOMAIN; 2.
 KW Phosphorylation; Repeat.
 FT DOMAIN 4 101
 FT PH 1
 FT DOMAIN 136 221
 FT DEP 2
 FT PH 2
 FT CONFLICT 120 120 L -> Q (IN REF. 3).
 FT CONFLICT 225 225 F -> L (IN REF. 3).
 SQ SEQUENCE 350 AA; 39900 MW; 348F3CB469B7CC53 CRC64;
 Query Match 75.0%; Score 33; DB 1; Length 350;
 Best Local Similarity 55.6%; Pred. No. 61;
 Matches 5; Conservative 3; Mismatches 1; Indels 0; Gaps 0;
 QY 1 LLGC:ITSL 9

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Db 292 LRGCVISV 300
! ! ! ! !
RESULT 9
ARGD_PSEPK STANFORD: PRT: 406 AA.
AC P59319;
DT 28-FEB-2003 (Rel. 41, Last sequence update)
DT 28-FEB-2003 (Rel. 42, Last annotation update)
DE Acetylornithine aminotransferase (EC 2.6.1.11) (NCOR1).
GN ARGD OR PP4481.
OS Pseudomonas putida (strain KT2440).
OC Bacteria; Proteobacteria; Gammaproteobacteria; Pseudomonadales;
OC Pseudomonadaceae; Pseudomonas.
OX NCBI_TaxID=160488;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=22423060; PubMed=12534463;
RA Nelson K.E., Weinel C., Paulsen I.T., Dodson R.J., Hilbert H.,
RA Martins dos Santos V.A.P., Feuts D.E., Gill S.R., Pop M., Holmes M.,
RA Brinkac C., Beanan M., Decoy R.T., Daugherty S., Koonsey J.,
RA Madupu R., Nelson W., White O., Peterson J., Khouri H., Hanco I.,
RA Chris Lee P., Holtzapple E., Scanlan D., Tran K., Motzke A.,
RA Utterback T., Rizzo M., Lee K., Kosack D., Moestl D., Wedler H.,
RA Lauber J., Stjepandic D., Hebeisel J., Straetz M., Heim S.,
RA Kiewitz C., Eisen J., Tirmis K.N., Duesterhoeft A., Truemmer B.,
RA Fraser C.W.;
RI "Complete genome sequence and comparative analysis of the
RI metabolically versatile Pseudomonas putida KT2440."
RL Environ. Microbiol. 4:799-808(2002).
CC -1- CATALYTIC ACTIVITY: N(2)-acetyl-L-ornithine + 2-oxoglutarate ->
CC acetyl-L-glutamate + 5-semialdehyde + 5-phosphate.
CC -1- COFACTOR: Pyridoxal phosphate (by similarity).
CC -1- PATHWAY: Arginine biosynthesis; fourth step.
CC -1- MISCELLANEOUS: May also have succinyl-L-aminopimelate
CC aminotransferase activity, thus carrying out the fourth step in
CC lysine biosynthesis.
CC -1- SIMILARITY: Belongs to class-II of pyridoxal-phosphate-dependent
CC aminotransferases.
CC
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CC or send an email to license@isb-sib.ch).
CC
CC EMBL: AF031898; AAC08059.1; -
CC InterPro: IPR003593; AAA_A-Pase.
CC DR Pfam: PF00448; SRP54.
CC DR Pfam: PF02381; SRP54.N.1.
CC DR Pfam: PF000819; SRP54.1.
CC DR SMART: SM00382; AAA.1.
CC DR PROSITE: PS00300; SRP54; FALSE_NEG.
CC DR FLAGELLIN: GTP-binding.
CC KW NP_BIND 225 232 GTP (BY SIMILARITY).
CC FT NP_BIND 303 307 GTP (BY SIMILARITY).
CC FT NP_BIND 362 364 GTP (BY SIMILARITY).
CC SQ SEQUENCE 406 AA; 43490 MW; 5597D2F10C3E221E CRC64;
Query Match 75.0%; Score 33; DB 1; Length 406;
Best Local Similarity 71.4%; Pred. No. 72;
Matches 5; Conservative 2; Mismatches 0; Indels 0; Gaps 0;
QY 1 LLGCVIT 7
Db 341 LRGCVIT 347
! ! ! ! !
RESULT 10
FLHF_PSEPU STANDARD: PRT: 437 AA.
ID FLHF_PSEPU
```

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AC 052256;
DT 15-DEC-1998 (Rel. 37, Created)
DT 15-DEC-1998 (Rel. 37, Last sequence update)
DT 15-DEC-1998 (Rel. 37, Last annotation update)
DE Flagellar biosynthesis protein flhF (Flagella associated GTP-binding
DE protein).
GN FLHF.
OS Pseudomonas putida.
OC Bacteria; Proteobacteria; Gammaproteobacteria; Pseudomonadales;
OC Pseudomonadaceae; Pseudomonas.
OX NCBI_TaxID=303;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=98164369; PubMed=9503621;
RA Ditty J.L., Grimm A.C., Harwood C.S.;
RA Identification of a chemotaxis gene region from Pseudomonas putida.;
RA FEMS Microbiol. Lett. 159:267-273(1998).
CC -1- FUNCTION: NECESSARY FOR FLAGELLAR BIOSYNTHESIS. MAY BE INVOLVED
CC IN TRANSLOCATION OF THE FLAGELLUM (BY SIMILARITY).
CC -1- SIMILARITY: BELONGS TO THE SRP FAMILY OF GTP-BINDING PROTEINS.
CC
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CC
CC EMBL: AF031898; AAC08059.1; -
CC InterPro: IPR003593; AAA_A-Pase.
CC DR Pfam: PF00448; SRP54.
CC DR Pfam: PF02381; SRP54.N.1.
CC DR Pfam: PF000819; SRP54.1.
CC DR SMART: SM00382; AAA.1.
CC DR PROSITE: PS00300; SRP54; FALSE_NEG.
CC DR FLAGELLIN: GTP-binding.
CC KW NP_BIND 225 232 GTP (BY SIMILARITY).
CC FT NP_BIND 303 307 GTP (BY SIMILARITY).
CC FT NP_BIND 362 364 GTP (BY SIMILARITY).
CC SQ SEQUENCE 437 AA; 47512 MW; EBFA6A650B9B27A CRC64;
Query Match 75.0%; Score 33; DB 1; Length 437;
Best Local Similarity 66.7%; Pred. No. 76;
Matches 6; Conservative 1; Mismatches 2; Indels 0; Gaps 0;
QY 1 LLGCVITSL 9
Db 355 LAGCVITSL 363
! ! ! ! !
RESULT 11
YBSE_PACSU STANDARD: PRT: 441 AA.
ID YBSE_PACSU
AC 005213; O08069; Q45578;
DT 28-FEB-2003 (Rel. 41, Created)
DT 28-FEB-2003 (Rel. 41, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Hypothetical protein ybde precursor.
GN ybde.
OS Bacillus subtilis.
OC Bacteria; Firmicutes; Bacillales; Bacillaceae; Bacillus.
OX NCBI_TaxID=1423;
RN [1]
RP SEQUENCE FROM N.A.
RX STRAIN=168;
RA MEDLINE=97419514; PubMed=9274029;
RA Liu H., Hage K., Yasumoto K., Ohashi Y., Yoshikawa H., Takahashi H.;
RA "Sequence and analysis of a 31 kb segment of the Bacillus subtilis
RA chromosome in the area of the trnH and trnG operons."
RA Microbiology 143:2763-2767(1997).
RL
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Db          591 MLCQVIT 597
|||||
RESULT 13
PGDS_HUMAN  STANDARD:      PRT: 1089 AA.
AC  P16234;
DI  01-APR-1390 (Rel. 14, Last sequence update)
DT  01-APR-1390 (Rel. 14, Last sequence update)
DT  15-SEP-2003 (Rel. 42, Last annotation update)
DE  Alpha platelet-derived growth factor receptor precursor (EC 2.7.1.112)
DE  (PDGF-R-alpha) (CD140a antigen).
CN  PDGFRA.
OS  Homo sapiens (Human).
OC  Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC  Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
CX  NCBI_TaxID=9606;
RN  [1]
RP  MEDLINE=89130149; PubMed=2536956;
RA  Matsui T., Heidaran M., Miki T., Popescu N., la Rochelle W.,
RA  Kraus M., Pierce J., Aaronson S.;
RT  Isolation of a novel receptor cDNA establishes the existence of two
RT  PDGF receptor genes.;
RL  Science 243:800-804(1989).
RN  [2]
RP  MEDLINE=89296915; PubMed=2544881;
RA  Claesson-Welsh L., Eriksson A., Westermark B., Hoidin G.H.;
RT  cDNA cloning and expression of the human A-type platelet-derived
RT  growth factor (PDGF) receptor establishes structural similarity to
RT  the B-type PDGF receptor.;
RL  Proc. Natl. Acad. Sci. U.S.A. 86:4917-4921(1989).
RN  [3]
RP  MEDLINE=95163874; PubMed=8585427;
RA  Kawagishi J., Ku T.;
RT  Structure, organization, and transcription units of the human
RT  alpha-platelet-derived growth factor receptor gene, PDGFRA.;
RL  Genomics 30:224-233(1995);
CC  -!- FUNCTION: THIS RECEPTOR BINDS PLATELET-DERIVED GROWTH FACTOR AND
CC  HAS A TYROSINE-PROTEIN KINASE ACTIVITY. THIS RECEPTOR CAN BIND
CC  EITHER PDGF-A OR PDGF-B.
CC  -!- CATALYTIC ACTIVITY: ATP + a protein:tyrosine - ADP + protein
CC  tyrosine phosphate.
CC  -!- SUBUNIT: DIMER OF EITHER ALPHA-ALPHA, BETA-BETA OR ALPHA-BETA
CC  SUBUNITS.
CC  -!- SUBCELLULAR LOCATION: Type I membrane protein.
CC  -!- SIMILARITY: BELONGS TO THE CSF-1/PDGF RECEPTOR FAMILY OF TYROSINE-
CC  PROTEIN KINASES.
CC  -!- SIMILARITY: Contains 5 immunoglobulin-like c2-type domains.
CC  -----
CC  THIS SWISS-PROT entry is copyright. It is produced through a collaboration
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DR  EMBL: D50010; BAA08742.1; JOINED.
DR  EMBL: D50011; BAA08742.1; JOINED.
DR  EMBL: D50012; BAA08742.1; JOINED.
DR  EMBL: D50013; BAA08742.1; JOINED.
DR  EMBL: D50014; BAA08742.1; JOINED.
DR  EMBL: D50015; BAA08742.1; JOINED.
DR  EMBL: D50016; BAA08742.1; JOINED.
DR  PIR: A40162; PFHUGA.
DR  HSP: P11362; IFGK.
DR  Genew; HGNC:8803; PDGFRA.
DR  MIM: 173490; -.
DR  GO: GO:0005887; C: integral to plasma membrane; TAS.
DR  GO: GO:0005018; F: platelet-derived growth factor, alpha-recp. .; TAS.
DR  GO: GO:0008283; P: cell proliferation; TAS.
DR  GO: GO:0007166; P: cell surface receptor linked signal transdu. .; TAS.
DR  InterPro: IPR007110; Ig-Like.
DR  InterPro: IPR003598; Ig_C2.
DR  InterPro: IPR003006; Ig_MHC.
DR  InterPro: IPR000719; Prot_Kinase.
DR  InterPro: IPR001324; RTKinase11.
DR  Pfam: PF00047; Ig_3.
DR  Pfam: PF00069; pkinase; 1.
DR  ProDom: PD000001; Prot_Kinase; 2.
DR  SMART: SM00408; IgC2; 1.
DR  SMART: SM00219; TyKc; 1.
DR  PROSITE: PS00835; IG_LIKE; 2.
DR  PROSITE: PS00107; PROTEIN_KINASE_ATP; 1.
DR  PROSITE: PS50011; PROTEIN_KINASE_DOM; 1.
DR  PROSITE: PS00109; PROTEIN_KINASE_TYR; 1.
DR  PROSITE: PS00240; RECEPTOR_TYR_KIN_III; 1.
KW  Tyrosine-protein kinase; Receptor; Transmembrane; Glycoprotein;
KW  Transferase; Phosphorylation; ATP-binding; Immunoglobulin domain;
KW  Signal; Repeat.
FT  SIGNAL 1 23
FT  CHAIN 24 1089
FT  DOMAIN 24 524
FT  TRANSMEM 525 549
FT  DOMAIN 550 1089
FT  DOMAIN 24 113
FT  DOMAIN 202 306
FT  DOMAIN 319 410
FT  DOMAIN 593 954
FT  DOMAIN 1041 1087
FT  NE_BIND 599 607
FT  BINDING 627 627
FT  ACT_SITE 818 818
FT  MOD_RES 849 849
FT  CARBOHYD 42 42
FT  CARBOHYD 76 76
FT  CARBOHYD 103 103
FT  CARBOHYD 179 179
FT  CARBOHYD 353 353
FT  CARBOHYD 359 359
FT  CARBOHYD 458 458
FT  CARBOHYD 468 468
SQ  SEQUENCE 1089 AA: 122669 MW: 5E3FB9940ACD1BE8 CRC64;
Query Match 75.0%; Score 33; DB 1; Length 1089;
Best Local Similarity 55.6%; Pred. No. 1,6e-02;
Matches 5; Conservative 3; Mismatches 1; Indels 0; Gaps 0;
QY 1 LIGCIITSL 9
DB 10 VLGLLTGL -8
RESULT 14
DICE_MOUSE
ID DICE_MOUSE STANDARD: PRT: 1906 AA.
AC Q8R418;
DT 28-FEB-2003. (Rel. 41, Created)
```


RA Nakajima C., Okazaki N., Yarakawa H., Kikuno A., Chara C., Nagase T.;
RT "Construction of expression-ready cDNA clones for KIAA genes: manual
RL curation of 330 KIAA cDNA clones.";
RL DNA Res. 9:99-106(2002).
RN [5]
RP SEQUENCE OF 1238-1912 FROM N.A.
RC TISSUE=Lung;
RX MEDLINE=99162526; PubMed=10051563;
RA Provost P., Samuelsson A., Radmark O.;
RI "Interaction of 5-lipoxygenase with cellular proteins.";
RL Proc. Natl. Acad. Sci. U.S.A. 96:1881-1885(1999).
CC -!- FUNCTION: Involved in cleaving double-stranded RNA in the RNA
CC interference (RNAi) pathway. It produces 21 to 23 bp dsRNAs
CC (siRNAs) which target the selective destruction of homologous
CC RNAs.
CC -!- SIMILARITY: BELONGS TO THE HELICASE FAMILY.
CC -!- SIMILARITY: Contains 2 RNase III domains.
CC -!- SIMILARITY: Contains 1 PAZ domain.
CC -!- SIMILARITY: Contains 1 DRRM (double-stranded RNA-binding) domain.
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CC
DR EMBL: AR028449; BAA78691.1; ALT_INIT.
DR EMBL: A313261; CAB38857.2; ..
DR EMBL: ABC23145; BAA76772.2; ALT_INIT.
DR Genew: HGNC:17098; DICER1.
DR MIM: 606241; ..
DR InterPro: IPR001410; DEAD.
DR InterPro: IPR001159; DS_RBD.
DR InterPro: IPR005034; DUF283.
DR InterPro: IPR001650; Helicase_C.
DR InterPro: IPR003100; PAZ.
DR InterPro: IPR005999; RNase_3.
DR Pfam: PF00270; DEAD_1.
DR Pfam: PF00035; GSRM; 1.
DR Pfam: PF03368; DUF283; 1.
DR Pfam: PF00271; Helicase_C; 1.
DR Pfam: PF02170; PAZ; 1.
DR Pfam: PF00636; Ribonuclease_3; 2.
DR SMART: SM00487; DEXDC; 1.
DR SMART: SM00358; DSRM; 1.
DR SMART: SM00490; HELIC_C; 1.
DR SMART: SM00535; RIBOC; 2.
DR PROSITE: PS0137; DS_RBD; 1.
DR PROSITE: PS0021; PAZ; 1.
DR PROSITE: PS00517; RNase_3; 1; 1.
DR PROSITE: PS0142; RNase_3; 2; 2.
KW Helicase; ATP-binding; RNA-binding; Hydrolyase; Nuclease; Endonuclease;
KW Repeat.
FT NP_BIND 14 41 ATP (POTENTIAL).
FT SITE 165 168 DECH_BOX.
FT DOMAIN 88 1632 PAZ.
FT DOMAIN 1266 1393 RNase III 1.
FT DOMAIN 1656 1814 RNase III 2.
FT DOMAIN 1839 1904 DRRM.
FT CONFLICT 65 80 VILTKELSYQIRGDFS -> STLLKSCLYLDAGETSA
FT CONFLICT 179 179 I -> F (IN REF. 1).
FT CONFLICT 185 185 N -> Z (IN REF. 1).
FT CONFLICT 204 204 C -> W (IN REF. 1).
FT CONFLICT 208 208 E -> D (IN REF. 1).
FT CONFLICT 213 213 I -> F (IN REF. 1).
FT CONFLICT 383 384 QQ -> HS (IN REF. 1).
FT CONFLICT 482 483 KQ -> NT (IN REF. 1).
FT CONFLICT 599 599 D -> H (IN REF. 1).
SQ SEQUENCE 1912 AA: 217627 MW: 996399DB4B074F21 CRC64:

Query Match 75.0%; Score 33; DR 1; Length 1912;
Best Local Similarity 75.0%; Pred. No. 2.6e+02;
Matches 6; Conservative 1; Mismatches 1; Indels 0; Gaps 0;
QY 1 LLGCITTS 8
Db 1556 LLGCYLTTS 1563
|||||
Search completed: September 29, 2003, 19:07:47
Job time : 13 secs

GenCore version 5.1.6
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OM protein - nucleic search, using frame_plus_p2c model

Run on: September 30, 2003, 10:34:19 : Search time 2022 Seconds
(without alignments)
167,226 Million cell updates/sec

Title: US-09-965-594-1_COPY_14_22
Perfect score: 44
Sequence: 1 LGGITSL 9

Scoring table: BLOSUM62
Xgapop 10.0 , Xgapext 0.5
Ygapop 10.0 , Ygapext 0.5
Fgapop 6.0 , Fgapext 7.0
Dgapop 6.0 , Delext 7.0

Searched: 266871 seqs, 2045481386 residues

Total number of hits satisfying chosen parameters: 5776890

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 99%
Listing first 45 summaries

Command line parameters:
-MODEL=frame_plus_p2c_model -DEV_X16
-Q/cqn211/USPIC_spool/US09965594/runat_23092003_164249_14072/app_query.fasta_1199
-DB=GenEmbl -QFMT=fastap -SUFFIX=rgc -MINMA=CH-0.1 -DOOP=0 -DOOPEXT=0
-UNITS=bits -START=1 -END=1 -MATRIX=blosum62 -TRANS=human40.cdi -LIST=45
-DOCALIGN=200 -START=1 -END=1 -THR_SCORE=pct -THR_MAX=99.9 -THR_MIN=0 -ALIGN=15 -MODE=LOCAL
-OUTFMT=ptc -NOR=ext -HEAPSIZE=13000 -MINLEN=0 -MAXLEN=200000000
-USER=US09965594 -CGN_1_13508 -runat_23092003_164249_14072 -NCPU=6 -ICPU=1
-NO_MMAP -LARGEQUERY -NGC_SCORES=0 -WAIT -DSPBCHECK=100 -LONGLOG
-DEV_TIMEOUT=120 -WARN_TIME=10 -THREADS=1 -XGAPOP=10 -XGAPEXT=0.5 -FGAPOP=6
-FGAPEXT=7 -YGAPOP=10 -YGAPEXT=0.5 -LFLOP=6 -DELEXT=7

Database : GenBank

1: gb_hum:
2: gb_hum:
3: gb_hum:
4: gb_hum:
5: gb_hum:
6: gb_hum:
7: gb_hum:
8: gb_hum:
9: gb_hum:
10: gb_hum:
11: gb_hum:
12: gb_hum:
13: gb_hum:
14: gb_hum:
15: gb_hum:
16: gb_hum:
17: gb_hum:
18: gb_hum:
19: gb_hum:
20: gb_hum:
21: gb_hum:
22: gb_hum:
23: gb_hum:
24: gb_hum:
25: gb_hum:
26: gb_hum:
27: gb_hum:
28: gb_hum:

29: em_vit:
30: em_hum:
31: em_hum:
32: em_hum:
33: em_hum:
34: em_hum:
35: em_hum:
36: em_hum:
37: em_hum:
38: em_hum:
39: em_hum:
40: em_hum:
41: em_hum:

pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	43	97.7	543	14	AB072047 Hepatitis
2	43	97.7	543	14	AB072048 Hepatitis
3	43	97.7	543	14	AB072067 Hepatitis
4	43	97.7	543	14	AB072102 Hepatitis
5	43	97.7	543	14	AB089533 Hepatitis
6	43	97.7	540	14	AB089539 Hepatitis
7	43	97.7	540	14	AB089558 Hepatitis
8	43	97.7	540	14	AB100809 Hepatitis
9	43	97.7	540	14	AB100827 Hepatitis
10	43	97.7	1211	14	AY070171 Hepatitis
11	43	97.7	2691	14	AF187824 Cache Val
12	43	97.7	2691	14	AF231116 Cache Val
13	43	97.7	2691	14	AF231118 Cache Val
14	43	97.7	4463	14	AF186241 Cache Val
15	43	97.7	9359	14	AF133016 Hepatitis
16	43	97.7	9359	14	AF165053 Hepatitis
17	43	97.7	9379	14	AF165049 Hepatitis
18	43	97.7	9379	14	AF165050 Hepatitis
19	43	97.7	9434	14	HPCJTB
20	43	97.7	9436	6	E07266 Blood-splea
21	43	97.7	9436	14	HPCJTB
22	43	97.7	9441	14	AY051292 Hepatitis
23	43	97.7	9598	14	AB049101 Hepatitis
24	43	97.7	217285	2	AC126317 Rattus no
25	43	97.7	260760	2	AC127861 Rattus no
26	42	95.5	543	14	AF169258 Hepatitis
27	42	95.5	819	6	E06893 DNA encodin
28	42	95.5	3461	6	E07544
29	42	95.5	3461	6	E09288
30	42	95.5	9375	14	AF207754 Hepatitis
31	42	95.5	9379	14	AF207768 Hepatitis
32	42	95.5	89352	9	AC006030 Homo sapi
33	42	95.5	105991	2	AC016960 Homo sapi
34	42	95.5	133801	3	AC034585 Homo sapi
35	42	95.5	138517	9	AC019288 Homo sapi
36	42	95.5	230396	2	AC103315 Rattus no
37	42	95.5	231275	2	AC099421 Rattus no
38	41	93.2	540	14	AB072056 Hepatitis
39	41	93.2	1096	14	AB013622 Hepatitis
40	41	93.2	8780	14	AF054253 Hepatitis
41	41	93.2	8780	14	AF054255 Hepatitis
42	41	93.2	8780	14	AF054259 Hepatitis
43	41	93.2	8781	14	AF054256 Hepatitis
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ALIGNMENTS

RESULT 1

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AB072047
LOCUS      AB072047          540 bp      RNA      linear      VRL 02-APR-2002
DEFINITION Hepatitis C virus type 1b gene for polyprotein, NS3 region, partial
            cds, isolate:H05-4.
ACCESSION AB072047
VERSION    AB072047.1  GI:19911838
KEYWORDS   Hepatitis C virus type 1b
SOURCE     Hepatitis C virus type 1b
            Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
            Hepacivirus.
REFERENCE  1
AUTHORS    Ogata,S.
TITLE      CORRELATION BETWEEN SECONDARY STRUCTURE OF AN AMINO-TERMINAL
            PORTION OF THE NONSTRUCTURAL PROTEIN 3 OF HEPATITIS C VIRUS AND
            DEVELOPMENT OF HEPATOCELLULAR CARCINOMA
JOURNAL    Unpublished
PUBLISHED  2 (bases 1 to 540)
REFERENCE  2
AUTHORS    Ogata,S.
TITLE      Direct Submission
JOURNAL    Submitted (26-Sep-2001) Satoshi Ogata, Kobe University, Graduate
            School of Medicine, Department of Microbiology, 7-5-1, Kusunoki-cho,
            Chuo-ku, Kobe, Hyogo 650-0017, Japan
            (E-mail:ogatasmed.kobe-u.ac.jp, Tel:81-78-382-5501,
            Fax:81-78-382-5519)
FEATURES   location/Qualifiers
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                        /cDS            <1..>540
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            BASE COUNT      DLYLVTRHADVIPVRRRGDGRGSLSPRPVSYLKSGSGGFLCPSGHVGVFRAAVCT
            ORIGIN          RGVAKAVDFVPVSMETMR"
                        1..>540
                        /gene="NS3"
                        /product="nonstructural protein"
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Percent Similarity: 100.00%      Conservative: 1
Best Local Similarity: 88.89%      Mismatches: 0
Query Match:    97.73%      Indels:      0
DB:             14      Gaps:      0
US-09-965-594-1_copy_14_22 (1-9) x AB072047 (1-540)
QY      1  LeuLeuglyCysIlelleThrSerIeu 5
            |||||.....|
Db      37 CTGCTGGTGCATGCTACTAGCCGC 63
RESULT  2
AB072048
LOCUS      AB072048          540 bp      RNA      linear      VRL 02-APR-2002
DEFINITION Hepatitis C virus type 1b gene for polyprotein, NS3 region, partial
            cds, isolate:H05-5.
ACCESSION AB072048
VERSION    AB072048.1  GI:19911840
KEYWORDS   Hepatitis C virus type 1b
SOURCE     Hepatitis C virus type 1b
            Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
            Hepacivirus.
REFERENCE  1
AUTHORS    Ogata,S.
TITLE      CORRELATION BETWEEN SECONDARY STRUCTURE OF AN AMINO-TERMINAL
            PORTION OF THE NONSTRUCTURAL PROTEIN 3 OF HEPATITIS C VIRUS AND
            DEVELOPMENT OF HEPATOCELLULAR CARCINOMA
JOURNAL    Unpublished
PUBLISHED  2 (bases 1 to 540)
REFERENCE  2
AUTHORS    Ogata,S.
TITLE      Direct Submission
JOURNAL    Submitted (26-Sep-2001) Satoshi Ogata, Kobe University, Graduate
            School of Medicine, Department of Microbiology, 7-5-1, Kusunoki-cho,
            Chuo-ku, Kobe, Hyogo 650-0017, Japan
            (E-mail:ogatasmed.kobe-u.ac.jp, Tel:81-78-382-5501,
            Fax:81-78-382-5519)
FEATURES   location/Qualifiers
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                        /gene="NS3"
                        /cDS            <1..>540
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                        /protein_id="BAB88230.1"
                        /db_xref="GI:19911841"
                        /translation="APITAYSQTRGLGCVITSLTGROKNOVEGEVQVSTATQSEFL
            mat_peptide      ATCVNGVGVTVFNGASKSLAGPKGPTQMTYINVDLDLVGKAPPGARSMPTCGSSL
            BASE COUNT      DLYLVTRHADVIPVRRRGDGRGSLSPRPVSYLKSGSGGFLCPSGHVGVFRAAVCT
            ORIGIN          RGVAKAVDFVPVSMETMR"
                        1..>540
                        /gene="NS3"
                        /product="nonstructural protein"
                        95 a 177 c 165 g 103 t
Alignment Scores:
Pred. No.:      4.14      Length:      540
Score:          43.00     Matches:      8
Percent Similarity: 100.00%      Conservative: 1
Best Local Similarity: 88.89%      Mismatches: 0
Query Match:    97.73%      Indels:      0
DB:             14      Gaps:      0
US-09-965-594-1_copy_14_22 (1-9) x AB072048 (1-540)
QY      1  LeuLeuglyCysIlelleThrSerIeu 9
            |||||.....|
Db      37 CTACTCGGTGCATGCTACTAGCCGC 63
RESULT  3
AB072067
LOCUS      AB072067          540 bp      RNA      linear      VRL 02-APR-2002
DEFINITION Hepatitis C virus type 1b gene for polyprotein, NS3 region, partial
            cds, isolate:L08.
ACCESSION AB072067
VERSION    AB072067.1  GI:19911878
KEYWORDS   Hepatitis C virus type 1b
SOURCE     Hepatitis C virus type 1b
            Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
            Hepacivirus.
REFERENCE  1
AUTHORS    Ogata,S.
TITLE      CORRELATION BETWEEN SECONDARY STRUCTURE OF AN AMINO-TERMINAL
            PORTION OF THE NONSTRUCTURAL PROTEIN 3 OF HEPATITIS C VIRUS AND
            DEVELOPMENT OF HEPATOCELLULAR CARCINOMA
JOURNAL    Unpublished
PUBLISHED  2 (bases 1 to 540)
REFERENCE  2
AUTHORS    Ogata,S.
TITLE      Direct Submission
JOURNAL    Submitted (26-Sep-2001) Satoshi Ogata, Kobe University, Graduate
            School of Medicine, Department of Microbiology, 7-5-1, Kusunoki-cho,
            Chuo-ku, Kobe, Hyogo 650-0017, Japan
            (E-mail:ogatasmed.kobe-u.ac.jp, Tel:81-78-382-5501,
            Fax:81-78-382-5519)
FEATURES   location/Qualifiers
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                        /cDS            <1..>540
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            BASE COUNT      DLYLVTRHADVIPVRRRGDGRGSLSPRPVSYLKSGSGGFLCPSGHVGVFRAAVCT
            ORIGIN          RGVAKAVDFVPVSMETMR"
                        1..>540
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                        /product="nonstructural protein"
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Alignment Scores:
Pred. No.:      4.14      Length:      540
Score:          43.00     Matches:      8
Percent Similarity: 100.00%      Conservative: 1
Best Local Similarity: 88.89%      Mismatches: 0
Query Match:    97.73%      Indels:      0
DB:             14      Gaps:      0
US-09-965-594-1_copy_14_22 (1-9) x AB072067 (1-540)
QY      1  LeuLeuglyCysIlelleThrSerIeu 9
            |||||.....|
Db      37 CTACTCGGTGCATGCTACTAGCCGC 63
RESULT  3
AB072067
LOCUS      AB072067          540 bp      RNA      linear      VRL 02-APR-2002
DEFINITION Hepatitis C virus type 1b gene for polyprotein, NS3 region, partial
            cds, isolate:L08.
ACCESSION AB072067
VERSION    AB072067.1  GI:19911878
KEYWORDS   Hepatitis C virus type 1b
SOURCE     Hepatitis C virus type 1b
            Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
            Hepacivirus.
REFERENCE  1
AUTHORS    Ogata,S.
TITLE      CORRELATION BETWEEN SECONDARY STRUCTURE OF AN AMINO-TERMINAL
            PORTION OF THE NONSTRUCTURAL PROTEIN 3 OF HEPATITIS C VIRUS AND
            DEVELOPMENT OF HEPATOCELLULAR CARCINOMA
JOURNAL    Unpublished
PUBLISHED  2 (bases 1 to 540)
REFERENCE  2
AUTHORS    Ogata,S.
TITLE      Direct Submission
JOURNAL    Submitted (26-Sep-2001) Satoshi Ogata, Kobe University, Graduate
            School of Medicine, Department of Microbiology, 7-5-1, Kusunoki-cho,
            Chuo-ku, Kobe, Hyogo 650-0017, Japan
            (E-mail:ogatasmed.kobe-u.ac.jp, Tel:81-78-382-5501,
            Fax:81-78-382-5519)
FEATURES   location/Qualifiers
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                        /cDS            <1..>540
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            BASE COUNT      DLYLVTRHADVIPVRRRGDGRGSLSPRPVSYLKSGSGGFLCPSGHVGVFRAAVCT
            ORIGIN          RGVAKAVDFVPVSMETMR"
                        1..>540
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                        /product="nonstructural protein"
                        95 a 177 c 165 g 103 t
Alignment Scores:
Pred. No.:      4.14      Length:      540
Score:          43.00     Matches:      8
Percent Similarity: 100.00%      Conservative: 1
Best Local Similarity: 88.89%      Mismatches: 0
Query Match:    97.73%      Indels:      0
DB:             14      Gaps:      0
US-09-965-594-1_copy_14_22 (1-9) x AB072067 (1-540)
QY      1  LeuLeuglyCysIlelleThrSerIeu 9
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Db      37 CTACTCGGTGCATGCTACTAGCCGC 63

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BASE COUNT      95 a   176 c   166 g   102 t
ORIGIN

Alignment Scores:
Pred. No.:      4.14      Length:      540
Score:          43.00      Matches:      8
Percent Similarity: 100.00%      Conservative: 1
Best Local Similarity: 88.89%      Mismatches: 0
Query Match:    97.73%      Indels:      0
DB:            14      Gaps:      0

US-09-965-594-1_COPY_14_22 (1-9) x AB089539 (1-540)

QY      1 LeuLeuGlyCysIleIleThrSerLeu 9
DB      37 CTACTTGGCTCGGTATCAGCAGCTC 63

RESULT 7
AB089558      Hepatitis C virus NS3 gene for polyprotein, partial cds,
LOCUS      AB089558      540 bp      RNA      linear      VRL 02-APR-2003
DEFINITION      Hepatitis C virus NS3 gene for polyprotein, partial cds,
ACCESSION      AB089558
VERSION      AB089558.1      GI:29467323
KEYWORDS
SOURCE      Hepatitis C virus
ORGANISM      Hepatitis C virus
VIRUSES; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
Hepadnavirus.

REFERENCE
1      Ogata,S. and Hotta,H.
AUTHORS      Identification of Hepatitis C Virus (HCV) Subtype 1b Strains That
TITLE      Are Highly, or Only Weakly, Associated with Hepatocellular
Carcinoma on the Basis of Secondary Structure of an Amino-Terminal
Portion of the HCV NS3 Protein
JOURNAL      Unpublished
REFERENCE      2      (bases 1 to 540)
AUTHORS      Hotta,H.
DIRECT SUBMISSION
JOURNAL      Submitted (07-AUG-2002) Hak Hotta, Kobe University Graduate School
of Medicine, Division of Microbiology; Chuo-ku, Kusunoki-cho 7-5-1,
Kobe, Hyogo 650-0017, Japan (E-mail:hotta@kobe-u.ac.jp,
Tel:81-78-382-5500, Fax:81-78-382-5519)
FEATURES
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location/Qualifiers
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DLYLTVRHADVPVRRSDGRGSLSPRVSYLKGSSGGLLCPGSHAVGIFRAAVCT
RGVAKAVDFVPVRESMETMR"
BASE COUNT      95 a   168 c   163 g   114 t
ORIGIN

Alignment Scores:
Pred. No.:      4.14      Length:      540
Score:          43.00      Matches:      8
Percent Similarity: 100.00%      Conservative: 1
Best Local Similarity: 88.89%      Mismatches: 0
Query Match:    97.73%      Indels:      0
DB:            14      Gaps:      0

US-09-965-594-1_COPY_14_22 (1-9) x AB089558 (1-540)

QY      1 LeuLeuGlyCysIleIleThrSerLeu 9

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DLYLTVRHADVPVRRSDGRGSLSPRVSYLKGSSGGLLCPGSHAVGIFRAAVCT
RGVAKAVDFVPVRESMETMR"
BASE COUNT      95 a   176 c   166 g   102 t
ORIGIN

Alignment Scores:
Pred. No.:      4.14      Length:      540
Score:          43.00      Matches:      8
Percent Similarity: 100.00%      Conservative: 1
Best Local Similarity: 88.89%      Mismatches: 0
Query Match:    97.73%      Indels:      0
DB:            14      Gaps:      0

US-09-965-594-1_COPY_14_22 (1-9) x AB089539 (1-540)

QY      1 LeuLeuGlyCysIleIleThrSerLeu 9
DB      37 CTACTTGGCTCGGTATCAGCAGCTC 63

RESULT 7
AB089558      Hepatitis C virus NS3 gene for polyprotein, partial cds,
LOCUS      AB089558      540 bp      RNA      linear      VRL 02-APR-2003
DEFINITION      Hepatitis C virus NS3 gene for polyprotein, partial cds,
ACCESSION      AB089558
VERSION      AB089558.1      GI:29467323
KEYWORDS
SOURCE      Hepatitis C virus
ORGANISM      Hepatitis C virus
VIRUSES; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
Hepadnavirus.

REFERENCE
1      Ogata,S. and Hotta,H.
AUTHORS      Identification of Hepatitis C Virus (HCV) Subtype 1b Strains That
TITLE      Are Highly, or Only Weakly, Associated with Hepatocellular
Carcinoma on the Basis of Secondary Structure of an Amino-Terminal
Portion of the HCV NS3 Protein
JOURNAL      Unpublished
REFERENCE      2      (bases 1 to 540)
AUTHORS      Hotta,H.
DIRECT SUBMISSION
JOURNAL      Submitted (07-AUG-2002) Hak Hotta, Kobe University Graduate School
of Medicine, Division of Microbiology; Chuo-ku, Kusunoki-cho 7-5-1,
Kobe, Hyogo 650-0017, Japan (E-mail:hotta@kobe-u.ac.jp,
Tel:81-78-382-5500, Fax:81-78-382-5519)
FEATURES
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RGVAKAVDFVPVRESMETMR"
BASE COUNT      95 a   168 c   163 g   114 t
ORIGIN

Alignment Scores:
Pred. No.:      4.14      Length:      540
Score:          43.00      Matches:      8
Percent Similarity: 100.00%      Conservative: 1
Best Local Similarity: 88.89%      Mismatches: 0
Query Match:    97.73%      Indels:      0
DB:            14      Gaps:      0

US-09-965-594-1_COPY_14_22 (1-9) x AB089558 (1-540)

QY      1 LeuLeuGlyCysIleIleThrSerLeu 9

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Db
|||||.....
37 CTACTGGTGCATCGCTCACTAGCCTC 63

RESULT 6
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LOCUS      Hepatitis C virus NS3 gene for polypeptide, partial cds, isolate:
DEFINITION Y-CH-06-B.
ACCESSION AB100809
VERSION    AB100809.1 GI:29467656
KEYWORDS
SOURCE
ORGANISM   Hepatitis C virus
            Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
            Hepacivirus.

REFERENCE
AUTHORS    Oyata,S. and Hotta,H.
TITLE       Identification of Hepatitis C Virus (HCV) Subtype 1b Strains That
            Are Highly, or Only Weakly, Associated with Hepatocellular
            Carcinoma on the Basis of Secondary Structure of an Amino-Terminal
            Portion of the HCV NS3 Protein
JOURNAL     Unpublished
PUBLISHED
AUTHORS     Hotta,H.
TITLE       Direct Submission
JOURNAL     Submitted (24-JAN-2003) Hak Hotta, Kobe University Graduate School
            of Medicine, Division of Microbiology; Chuo-ku, Kusunoki-cho 7-5-1,
            Kobe, Hyogo 650-0017, Japan (E-mail:hotta@kobe-u.ac.jp).
            Tel:81-78-382-5500, Fax:81-78-382-5513
            Location/Qualifiers
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            1..540
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                /db_xref="GI:29467657"
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                RGVAKAVDFEVEVSMETMR"
            BASE COUNT      93 a 171 c 166 g 110 t
            ORIGIN
            Alignment Scores:
            Pred. No.:      4.14      Length:      540
            Score:          43.00      Matches:      8
            Percent Similarity: 100.00%      Conservative: 1
            Best Local Similarity: 88.89%      Mismatches: 0
            Query Match:      97.73%      Indels:      0
            Db:              14      Gaps:      0
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QY      1 LeuGugGlyCysIleIeThrSerLeu 9
|||||.....
37 CTACTGGTGCATCGCTCACTAGCCTC 63

RESULT 10
AB100809
LOCUS      Hepatitis C virus strain 687 polyprotein gene, partial cds.
DEFINITION AY070171
ACCESSION  AY070171
VERSION     AY070171.1 GI:19879463
KEYWORDS
SOURCE      Hepatitis C virus
            Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
            Hepacivirus.
REFERENCE   1 (bases 1 to 1211)
AUTHORS     Kaimina,O., Norder,H., Mukomolov,S. and Magnus,L.O.
TITLE       A natural intergenotypic recombinant of hepatitis C virus
            identified in St. Petersburg
JOURNAL     J. Virol. 76 (8), 4034-4043 (2002)
MEDLINE     21904745
PUBMED      11907242
REFERENCE   2 (bases 1 to 1211)

```


AUTHORS Kalinina, G., Nordcr, H., Mukomolov, S. and Maynits, L.
 TITLE Direct Submission
 JOURNAL Submitted (12-DEC-2001) Virology, Swedish Institute for Infectious
 Disease Control, Solna SE 171 82, Sweden

FEATURES

1..1211
 Location/Qualifiers
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 /db_xref="GI:19879464"

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 VCFNPNNAVCTE:NAIEBK:TDIELOKTVTITPLNLAQNHKLYSGQINILGS
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BASE COUNT 225 a 358 c 349 g 279 t
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 Query Match: 97.73% Indels: 0
 DB: 14 Gaps: 0

US-09-965-594-1_COPY_14_22 (1-9) x AY070171 (1-1211)

Qy 1 LeuLeuGlyCysIleIleThrSerLeu 9
 Db 969 CTACTGGGCGCATGCTGACTAGCCTC 995

RESULT 11

AF187824/c
 LOCUS AF187824 2691 bp RNA linear VRL 02-MAY-2001
 DEFINITION Cache Valley virus 8063 GI gene, partial cds.
 ACCESSION AF187824
 VERSION AF187824.1 GI:6959495

KEYWORDS
 SOURCE Cache Valley virus
 ORGANISM Cache Valley virus

Viruses; ssRNA negative-strand viruses; Bunyaviridae;
 Orthobunyavirus; Bunyamwera virus group.

1 (bases 1 to 2691)
 Brockus, C.L. and Grimstad, P.R.

REFERENCE
 AUTHORS Comparative analysis of GI glycoprotein-coding sequences of Cache
 TITLE Valley virus (Bunyaviridae: Bunyavirus) isolates

JOURNAL Virus Genes 22 (2), 133-139 (2001)
 MEDLINE 21221452
 PUBMED 11324749

REFERENCE

2 (bases 1 to 2691)
 Brockus, C.L., Collins, F.H., Besansky, N.J., and Grimstad, P.R.

REFERENCE
 TITLE Phylogenetic Analysis of Cache Valley Virus Isolates

JOURNAL Unpublished
 3 (bases 1 to 2691)
 Brockus, C.L., and Grimstad, P.R.

REFERENCE

AUTHORS Direct Submission
 TITLE Submitted (17-SEP-1999) Vector Biology Labs, University of Notre
 JOURNAL Dame, Galvin Life Sciences Center, Notre Dame, IN 46556, USA

FEATURES

Location/Qualifiers
 1..2691
 /organism="Cache Valley virus"
 /mol_type="genomic RNA"

/strain="PC68"

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/country="USA: Dennisville, NJ"

/note="1982 isolate from Ae. sollicitans"

1..2691

/note="envelope glycoprotein; putative soluble portion"

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/product="GI"

/protein_id="AA533116.1"

/db_xref="GI:6959496"

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 CMLFYTFNAPGPTESFLYQLOKNTTGASKVLGLI:TKYGNMNFVGVWKEGQY
 LTLTYNNTLSGNRLILATQFALSRSHSGROPSMSNAIOGSLTKECHNAKGVGC
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 VNKRKELFLTDVSEVDIFSVAS:CKMADGVCIVNCAKNWIKDGLIYFTDHR
 REGQDNDGFEYCSISGNTERFP:NPDI:EKDCTWEPHSRKSXYISILSDSBEFR
 RAMAKISHTLIITNFKPTANLPH:RPTKYFTANGVNSDGI:ESAYLLSIPALGGI
 SDGVQSDSSEFLDMIVFDISAVITTYVHYDTGHTIGIY:OHDHCTGPPAMV
 PHKENWNTSOFRTSMGFEPCGLAINTGCVFGSCOD:THPETKYRKAEEVVLIT
 VCFNPNNAVCTE:NAIEBK:TDIELOKTVTITPLNLAQNHKLYSGQINILGS
 FSQCGNOKNINSILGKMTV:KFDYVCHGASKEQVIVRRYNNNFSCALLPEFTSLI
 FADNHTELVAKTHLDELQFKIM:GDIRYKSPAFSELEIDAKCVGCGPCFESTISC
 NFQVITNIDTVCQVEGECTLFHNRI:ISAMKSGYGLKMSCOTKPNONEFTICNRRYS
 VLETT:DKNDK:EVNTGDOTSYLYEKDSRCKTWLCRVDEG"

BASE COUNT 949 a 476 c 539 g 727 t
 ORIGIN

Alignment Scores:
 Pred. No.: 21.3 Length: 2691
 Score: 43.00 Matches: 8
 Percent Similarity: 100.00% Conservative: 1
 Best Local Similarity: 88.89% Mismatches: 0
 Query Match: 97.73% Indels: 0
 DB: 14 Gaps: 0

US-09-965-594-1_COPY_14_22 (1-9) x AF187824 (1-2691)

Qy 1 LeuLeuGlyCysIleIleThrSerLeu 9
 Db 1367 ITGTTAGGCTGTGATACCTTCTCTG 1361

RESULT 12

AF231116/c

LOCUS AF231116 2691 bp DNA linear VRL 02-MAY-2001

DEFINITION Cache Valley virus isolate Ar78-5912 GI envelope glycoprotein gene,
 partial cds.

ACCESSION AF231116

VERSION AF231116.1 GI:13249094

KEYWORDS
 SOURCE Cache Valley virus

ORGANISM Cache Valley virus

Viruses; ssRNA negative-strand viruses; Bunyaviridae;
 Orthobunyavirus; Bunyamwera virus group.

1 (bases 1 to 2691)
 Brockus, C.L. and Grimstad, P.R.

REFERENCE
 AUTHORS Comparative analysis of GI glycoprotein-coding sequences of Cache
 TITLE Valley virus (Bunyaviridae: Bunyavirus) isolates

JOURNAL Virus Genes 22 (2), 133-139 (2001)
 MEDLINE 21221452
 PUBMED 11324749

REFERENCE

2 (bases 1 to 2691)
 Brockus, C.L. and Grimstad, P.R.

REFERENCE
 TITLE Direct Submission

JOURNAL Submitted (04-FEB-2000) Biological Sciences, University of Notre
 Dame, Galvin Life Sciences Bldg, Notre Dame, IN 46556, USA

FEATURES

Location/Qualifiers
 1..2691
 /organism="Cache Valley virus"
 /mol_type="genomic DNA"
 /isolate="Ar78-5912"
 /specific_host="Anopheles quadrimaculatus"

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RAMAKUSHLLIYNKPTANPHIRPTTKFTANGVENSQIESAFLSSIPALGGI
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VCINFGNNYCTEINAEIKPTDEIELOQKVTDTKLPNLLAVONHKLKLYSGQINDLGS
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FADNHETLEVAHTKHLIGELQKIMLDIRYKFAESPELEIDAKCVGCSQESYSC
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BASE COUNT 947 a 475 c 541 g 728 t
ORIGIN

Alignment Scores:
Pred. No.: 21.3 Length: 2269
Score: 43.00 Matches: 8
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 88.89% Mismatches: 0
Query Match: 97.73% Indels: 0
DB: 14 Gaps: 0

US-09-965-594-1_COPY_14_22 (1-9) x AF231118 (1-2269)

QY 1 LeuLeuGlyCysIleIleThrSerLeu 9
|||||
DB 1387 TTGTTAGGTTGTGTGATACTTCCTC 1361

RESULT 13
AF231118/c
LOCUS
Cache Valley virus isolate 61D240 G1 envelope glycoprotein gene,
partial cds.
AF231118
AF231118.1 G1:13249096
Cache Valley virus
Cache Valley virus
Viruses: ssRNA negative-strand viruses; Bunyaviridae;
Orthobunyavirus; Bunyamwera virus group.
1 (bases 1 to 2269)
Brookus,C.L. and Grimstad,P.R.
Comparative analysis of G1 glycoprotein-coding sequences of Cache
Valley virus (Bunyaviridae: Bunyavirus) isolates
Virus Genes 22 (2), 133-139 (2001)
JOURNAL
MEDLINE
2122452
PUBMED
2 (bases 1 to 269)
Brookus,C.L. and Grimstad,P.R.
Direct Submission
AUTHORS
Submitted (04-FEB-2003) Biological Sciences, University of Notre
Dame, Galvin Life Sciences Bldg, Notre Dame, IN 46556, USA
JOURNAL
FEATURES
source
1..269;
/organism="Cache Valley virus"
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/isolate="61D240"
/specific_host="Psorophora cingulata"
/db_xref="taxon:80935"
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/note="isolated in 1961"

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VNKIKLTCFLIDVSVVDVFSVAASPCMKADKGVCTVNDKWNIIKCSGLYFTDH
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VCINFGNNYCTEINAEIKPTDEIELOQKVTDTKLPNLLAVONHKLKLYSGQINDLGS
VSCGCGNVQKTNHSLGMGTAKFDYVCHGASRKDVIVRCYNNKDFSCLLKEETSLI
FADNHETLEVAHTKHLIGELQKIMLDIRYKFAESPELEIDAKCVGCSQESYSC
NFQIVNTDVCVSEGPCFLFHNRIISANKQSYGLKMSCQTKPNONEEFFICNRKYS
VFETIDKNDKIEVNTGDTQTSYIYEKDSRCKTMCWRVDEG"
BASE COUNT 962 a 468 c 535 g 726 t
ORIGIN

Alignment Scores:
Pred. No.: 21.3 Length: 2269
Score: 43.00 Matches: 8
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 88.89% Mismatches: 0
Query Match: 97.73% Indels: 0
DB: 14 Gaps: 0

US-09-965-594-1_COPY_14_22 (1-9) x AF231118 (1-2269)

QY 1 LeuLeuGlyCysIleIleThrSerLeu 9
|||||
DB 1387 TTGTTAGGTTGTGTGATACTTCCTC 1361

RESULT 14
AF186241/c
LOCUS
Cache Valley virus strain: M180-1-450 M segment, complete sequence.
AF186241
AF186241.1 G1:659489
Cache Valley virus
Cache Valley virus
Viruses: ssRNA negative-strand viruses; Bunyaviridae;
Orthobunyavirus; Bunyamwera virus group.
1 (bases 1 to 4463)
Brookus,C.L. and Grimstad,P.R.
Comparative analysis of G1 glycoprotein-coding sequences of Cache
Valley virus (Bunyaviridae: Bunyavirus) isolates
Virus Genes 22 (2), 133-139 (2001)
JOURNAL
MEDLINE
2122452
PUBMED
2 (bases 1 to 4463)
Brookus,C.L., Collins,P.H., Besanaky,N.L. and Grimstad,P.R.
Phylogenetic analysis of Cache Valley virus isolates
Unpublished
3 (bases 1 to 4463)
Brookus,C.L. and Grimstad,P.R.
Direct Submission
AUTHORS
Submitted (14-SEP-1999) Vector Biology Labs, University of Notre
Dame, Galvin Life Sciences Center, Notre Dame, IN 46556, USA
JOURNAL
FEATURES
source
1..4463
/organism="Cache Valley virus"
/mol_type="genomic RNA"
/strain="M180-1-450"
/db_xref="taxon:80935"
/chromosome="M segment"

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 GNIMVAPILKAMITHEFFSILLAGEKLEKALDCQYACYSLEPLDLPQLIQKH
 GLSAPSIHSYSGEINRVAACLEKLGVTPLVANHRAVRAKLSQGRANCKYIL
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 CIVILITNE*

BASE COUNT 1877 a 2830 c 2680 g 1468 t 4 others
 ORIGIN

Alignment Scores:
 Pred. No.: 76 Length: 9359
 Score: 43.00 Matches: 8
 Percent Similarity: 100.00% Conservative: 1
 Best Local Similarity: 88.89% Mismatches: 0
 Query Match: 97.73% Indels: 0
 DB: 14 Gaps: 0

US-09-965-594-1_COPY_14_22 (1-9) x AF113916 (1-9359)

Oy 1 LeuLeuGlyCysIleIleThrSerLeu 9
 :|||||:|||||:|||||:|||||:|||||
 Db 3444 CTACTTGGCTSCATCG:CAACAGCCIC 3470

Search completed: September 30, 2003, 12:51:59
 Job time : 2207 secs

SanCore version 5.1.6
Copyright (c) 1993 - 2003 Computer Aid

OM protein - nucleic search, using frame_plus_pdn model

Run on: September 29, 2003, 19:05:03 : Search time 139 Seconds
(without alignments)
128,545 Million cell updates/sec

Title: us-09-965-594-1_copy_14_22
Perfect score: 44
Sequence: 1 LISC1:TSL 9

Scoring table: R-OSUM62
Xgapop 10.0 , Xgapext 0.5
Ygapop 10.0 , Ygapext 0.5
Fgapop 6.0 , Fgapext 7.0
Delop 6.0 , Delext 7.0

Searched: 2552756 seqs, 1349719017 residues

Total number of hits satisfying chosen parameters: 5105271

Minimum DB seq length: 0
Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%
Maximum Match 99%
Listing first 45 summaries

Command line parameters:
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-O=Cg22_1/USPTO_spoos/US09565594/runat_29032003_154249_14063/app_query.fasta.1.199
-DB=N_Geneseq_19Jun03 -QMT=fastap -SUFFIX=ring -MINNA=CH=0.1 -LOOPEXT=0
-LOOPEXT=0 -UNITS=bits -START=1 -END=1 -MATRIX=BLOCK62 -TRANS=human40.cdi
-LIST=45 -LOCAL=LOCAL -OUTFMT=ptc -THR_SCORE=ptc -THR_MAX=99.9 -THR_MIN=0 -ALIGN=15
-MODE=LOCAL -OUTFMT=ptc -NORM=score -HEAPSIZE=500 -MINLEN=0 -MAXLEN=200000000
-USER=US09565594 -SCGN=1.1.312 -runat_29032003_164247_14064 -NCPU=6 -ICPU=3
-NO_MMAP -LARGEQUERY -NEG_SCORES=0 -WAIT -DSP510CK=100 -LONG3CG
-DEV_TIMEOUT=220 -WARN_TIMEOUT=30 -THREADS=1 -XGAPOP=10 -XGAPEXT=0.5 -FSAPOP=6
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Database : N_Geneseq_19Jun03:
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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed.

and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	43	97.7	9436	15	AA063499 Blood transmissible
2	42	95.5	819	15	AA058517 NANBH virus gene f
3	42	95.5	3461	15	AA064368 Non-A, non-B hepat
4	42	95.5	3461	15	AA064368 5'UTR/CORE/ESV/NS1
5	41	93.2	4977	23	AA058668 DNA encoding novel
6	41	93.2	4977	23	AA058668 DNA encoding novel
7	41	93.2	4977	23	AA058668 DNA encoding novel
8	41	93.2	9595	20	AA058668 Infectious hepatitis
9	41	93.2	9595	22	AA058668 Nucleotide sequenc
10	41	93.2	9595	22	AA058668 Infectious Hepatit
11	41	93.2	94895	21	AA058668 BAC containing rep
12	41	93.2	94895	21	AA058668 BAC containing rep
13	41	93.2	103929	21	AA058668 BAC containing rep
14	41	93.2	134499	21	AA058668 BAC containing rep
15	41	93.2	1082138	21	AA058668 Arabidopsis thalia
16	41	93.2	1082138	21	AA058668 Arabidopsis thalia
17	40	90.9	424	24	ABL63442 Breast cancer rela
18	40	90.9	1933	20	AA058668 HCV NS3 DNA. Hepa
19	40	90.9	2346	24	AA058668 cDNA encoding huma
20	40	90.9	2520	25	AA058668 Human novel polynu
21	40	90.9	8145	20	AA058668 Plasmid pBT-BS(+)
22	40	90.9	910715	20	AA058668 Porreilia purgoorie
23	39	88.6	4154	25	AA058668 Aspergillus fumiga
24	39	88.6	4746	21	AA058668 Rat calcium/calmod
25	39	88.6	9402	14	AA058668 Human Hepatitis C
26	39	86.4	9405	14	AA058668 Full-length Hepati
27	38	86.4	242	25	AA058668 Human GDP-mannose
28	37	84.1	39	20	AA058668 HCV NS4A-NS3 compl
29	37	84.1	39	20	AA058668 HCV NS4A-NS3 compl
30	37	84.1	39	20	AA058668 HCV NS4A-NS3 compl
31	37	84.1	39	20	AA058668 HCV NS4A-NS3 compl
32	37	84.1	39	20	AA058668 HCV NS4A-NS3 compl
33	37	84.1	39	20	AA058668 HCV NS4A-NS3 compl
34	37	84.1	39	20	AA058668 HCV NS4A-NS3 compl
35	37	84.1	39	20	AA058668 HCV NS4A-NS3 compl
36	37	84.1	39	20	AA058668 HCV NS4A-NS3 compl
37	37	84.1	39	20	AA058668 HCV NS4A-NS3 compl
38	37	84.1	39	20	AA058668 HCV NS4A-NS3 compl
39	37	84.1	39	20	AA058668 HCV NS4A-NS3 compl
40	37	84.1	39	20	AA058668 HCV NS4A-NS3 compl
41	37	84.1	39	20	AA058668 HCV NS4A-NS3 compl
42	37	84.1	39	20	AA058668 HCV NS4A-NS3 compl
43	37	84.1	39	20	AA058668 HCV NS4A-NS3 compl
44	37	84.1	39	20	AA058668 HCV NS4A-NS3 compl
45	37	84.1	39	20	AA058668 HCV NS4A-NS3 compl

ALIGNMENTS

RESULT 1
AA063499
CD AA063499 standard: cDNA; 9435 BP.
XX AA063499;
AC AA063499;
XX
XX 17-JAN-1995 (first entry)

DE Flood transmissible NANBHV genome.

XX Polymerase chain reaction; PCR; amplify; primer; non-A, non-B hepatitis;
KW NANBH; virus; blood transmissible; detection; hepatitis virus; RT-PCR;
KW C100 antibody; HCV RNA; NS5 region; ds.
XX Non-A, non-B hepatitis virus.

XX
XX Key Location/Qualifiers
FH

```

FT CDS 342..9374
FT /*tag= a
FN JP06105690-A.
PD 19-APR-1994.
XX
XX 10-MAR-1992; 92JP-0051885.
XX 10-MAR-1992; 92JP-0051885.
XX (KAEN/) KAENNO K.
PA
DR WPI; 1994-163139/20.
DR P-PSDB; AAR53417.
XX
XX Blood-transmissible non-A non-B hepatitis virus DNA - used for
PT detection of hepatitis virus
PT
PS Claim 1; Page 8-20; 22pp; Japanese.
XX
XX This sequence represents the genome of a blood transmissible non-A.
CC non-B hepatitis (NANBH) virus. This sequence was isolated using the
CC primers given in AAQ63500-95. The amplified fragments are used in the
CC detection of hepatitis virus. This target DNA was isolated from serum
CC of chronically infected NANBH patients who were C100 antibody-positive
CC and HCV RNA (NS5 region) positive. Reverse transcription-PCR and PCR
CC were performed on cDNA and the total human NANBH DNA was constructed
CC from 23 clones.
XX
SQ Sequence 9436 BP; 1876 A; 2840 C; 2055 G; 1974 T; 91 other;

Alignment Scores:
Pred. No.: 653 Length: 9436
Score: 43.00 Matches: 8
Percent Similarity: 100.00% Conservativeness: 1
Best Local Similarity: 88.89% Mismatches: 0
Query Match: 97.73% Indels: 0
DB: 15 Gaps: 0

US-09-965-594-1_COPY_14_22 (1-9) x AAQ63459 (1-9436);

RESULT 2
AAQ58617
ID AAQ58817 standard; cDNA: 819 BP.
XX
XX AAQ58817;
DI 24-NOV-1994 (first entry)
DE NANBH virus gene fragment #4.
XX
XX Antigen; structural; non-structural; non A non B hepatitis virus;
KW NANBH; NANBH; patient; plasma; diagnosis; detection; carrier; ss.
XX
XX Non A Non B hepatitis virus.
XX
XX JP06070778-A.
XX
XX 15-MAR-1994.
XX
XX 01-JUN-1993; 92JP-0156097.
XX
XX 10-JUL-1992; 92JP-0207391.
XX
XX (KOKU-) KOKUSAI SHIYAKU KK.
PA (SANW ) SANWA KAGAKU KENKYUSHO CO.
PA (TOFU ) TONEN CORP.
PA (TOKR-) ZH TOKYOOTO RINSHO IGAKU SOGO KENKYUSHO.

```

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XX WPI; 1994-128677/16.
XX P-PSDB; AAR50072.
XX
XX Nucleic acid fragment coding non-A non-B hepatitis virus antigen
PT - useful in diagnosis of NANB patient and detection of virus
PT carrier
XX
XX Claim 8; Page 18-19; 37pp; Japanese.
XX
XX The sequences given in AAQ58814-27 encode antigens of structural and
CC non-structural regions of non A non B hepatitis virus (NANBHV).
CC These sequences were derived from the plasma of a NANBH patient by
CC recombinant DNA techniques. These fragments are useful for the
CC diagnosis of NANBH patients and the detection of NANBH carriers.
XX
XX Sequence 819 BP; 150 A; 255 C; 241 G; 173 T; 0 other;

Alignment Scores:
Pred. No.: 664 Length: 819
Score: 42.00 Matches: 8
Percent Similarity: 100.00% Conservativeness: 1
Best Local Similarity: 88.89% Mismatches: 0
Query Match: 95.45% Indels: 0
DB: 15 Gaps: 0

US-09-965-594-1_COPY_14_22 (1-9) x AAQ58817 (1-819)

QY 1 LeuLeuGlyCysIleIleThrSerLeu 9
IIIIIIIIIIIIIIIIIIIIII
DB 67 CTAATGCTGCATCATCAGCAGCTC 93

RESULT 3
AAQ64068
ID AAQ64068 standard; cDNA: 3461 BP.
XX
XX AAQ64068;
DI 14-FEB-1995 (first entry)
DE Non-A, non-B hepatitis virus gene #4.
XX
XX Non-A, non-B hepatitis virus; NANBHV; hepatitis C virus; HCV;
KW core; ENV; NS1; NS2; NS3; antigen; detection; ss.
XX
XX Hepatitis C virus.
XX
XX Key Location/Qualifiers
FT CDS 307..3461
FT /*tag= a
FT misc_RNA 307..879
FT /*tag= b
FT misc_RNA 880..1455
FT /*tag= c
FT misc_RNA 1456..2736
FT /*tag= d
FT misc_RNA 2737..3461
FT /*tag= e
FT /*label= NS2-3
FT /*note= "NS3 N-terminal"
XX
XX JP06141870-A.
XX
XX 24-MAY-1994.
XX
XX 12-MAR-1992; 92JP-0088140.
XX
XX 12-MAR-1992; 92JP-0088140.
XX (SANW ) SANWA KAGAKU KENKYUSHO CO.
PA

```

PA (TOFU) TONEN CORP.
PA (TOKR-) 2H TOKYOTO RINSHO IGAKU SOG KENKYUSEI.
XX
DR WPI: 1994-22026/25.
DR P-PSDB: AAR54066.
XX
XX DNA coding a Non-A, non-B hepatitis virus antigen - useful for
PT detecting HCV within serum
PT
XX
XX Claim 1-5; Page 11-15; 22pp; Japanese.
PS
XX Hepatitis C virus #4 and #6 genes were isolated (AA064068-69).
CC Both genes contain the core, EN1, NS1, NS2 and NS3 regions.
CC A core region fragment is given in AA064067.
XX
XX Sequence 3461 BP; 638 A; 1046 C; 1012 G; 765 T; 0 other;
SQ

Alignment Scores:
Pred. No.: 336 Length: 3461
Score: 42.00 Matches: 8
Percent Similarity: 100.00% Conservative: 1
Best Local Similarity: 88.89% Mismatches: 0
Query Match: 95.45% Indels: 0
DB: 15 Gaps: 0

US-09-965-594-1_COPY_14_22 (1-9) x AA064068 (1-3461)

QY 1 LeuLeuGlyCysIleIleThrSerLeu 9
ID AAT130386
DB 3421 CTAATTGGTGCATCATCACCAGCCTA 3447

RESULT 4
AA030386
ID AAT130386 standard: cDNA: 3461 BP.
XX
AC AAT130386;
XX
XX 22-AUG-1996 (first entry)
XX
DE 5'UTR/CORE/ENV/NS1/NS2/NS3 cDNA from HCV (#4).
XX
XX Hepatitis C virus; HCV; antigen; detection; antibody; ds.
XX
OS Hepatitis C virus.
XX
FH Key Location/Qualifiers
FI CDS 307..3461
FT /*tag= a
FT misc_feature 307..373
FT /*tag= b
FT /*product= core peptide
FT misc_feature 880..1453
FT /*tag= c
FT /*product= ENV;
FT misc_feature 1456..2736
FT /*tag= d
FT /*product= NS1/ENV2
FT misc_feature 2737..3461
FT /*tag= e
FT /*product= NS2 and NS3
XX
XX JP07133291-A.
PN
XX
XX 23-MAY-1995.
PC
XX
XX 18-JUN-1993; 93JP-0147944.
PF
XX
XX 16-JUN-1993; 93JP-0147944.
PR
XX (TOFU) TONEN CORP.
PA
XX
XX WPI: 1995-220780/29.
DR
DR P-PSDB: AAR98361.

XX Recombinant polypeptide comprising partial NS1 region of hepatitis
PT non-A non-B viral antigen - used in a method for detecting
PT antibodies against hepatitis non-A non-B virus.
XX
XX Disclosure: Page 10-12; 15pp; Japanese.
XX
XX The sequences given in AAT30386-27 encode the 5'UTR/CORE/ENV/NS1/NS2/
CC NS3 protein region derived from hepatitis C virus (HCV) isolates #4
CC and #6 respectively. The proteins encoded by these sequences partic-
CC encode amino acids 394-495 of the HCV NS1 antigen. These protein
CC fragments may be used in the detection of antibodies against HCV.
XX
XX Sequence 3461 BP; 638 A; 1046 C; 1012 G; 765 T; 0 other;
SQ

Alignment Scores:
Pred. No.: 336 Length: 3461
Score: 42.00 Matches: 8
Percent Similarity: 100.00% Conservative: 1
Best Local Similarity: 88.89% Mismatches: 0
Query Match: 95.45% Indels: 0
DB: 16 Gaps: 0

US-09-965-594-1_COPY_14_22 (1-9) x AAT30386 (1-3461)

QY 1 LeuLeuGlyCysIleIleThrSerLeu 9
ID AAT30386
DB 3421 CTAATTGGTGCATCATCACCAGCCTA 3447

RESULT 5
AAS86668/C
ID AAS86668 standard: cDNA: 4977 BP.
XX
AC AAS86668;
XX
XX 13-FEB-2002 (first entry)
XX
DE DNA encoding novel human diagnostic protein #22472.
XX
KW Human; chromosome mapping; gene mapping; gene therapy; forensic;
KW food supplement; medical imaging; diagnostic; genetic disorder; ss.
XX
XX Homo sapiens.
XX
XX NC000175067-A2.
PN
XX 11-OCT-2001.
PO
XX
XX 30-MAR-2001; 2001NC-0508631.
PF
XX
XX 31-MAR-2000; 2000NC-0540217.
PR
XX 23-AUG-2000; 2000US-0649167.
XX
XX (HYSEP-) HYSEQ INC.
XX
XX DimaMac RT, Liu C, Tang YT;
XX
XX WPI: 2001-039362/73.
DR
DR P-PSDB: ABC22481.
XX
XX Now isolated polynucleotide and encoded polypeptides, useful in
PT diagnostics, forensics, gene mapping, identification of mutations
PT responsible for genetic disorders or other traits and to assess
PT biodiversity -
XX
XX Claim 1: SEQ ID NO 22472; 103pp; English.
PS
XX
XX The invention relates to isolated polynucleotide (I) and
CC polypeptide (II) sequences. (I) is useful as hybridisation probes,
CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome
CC and gene mapping, and in recombinant production of (II). The
CC polynucleotides are also used in diagnostics as expressed sequence tags
CC for identifying expressed genes. (I) is useful in gene therapy techniques

CC to restore normal activity of (II) or to treat disease states involving
 CC (II). (II) is useful for generating antibodies against it, detecting or
 CC quantitating a polypeptide in tissue, as molecular weight markers and as
 CC a food supplement. (II) and its binding partners are useful in medical
 CC imaging of sites expressing (II). (I) and (II) are useful for treating
 CC disorders involving aberrant protein expression or biological activity.
 CC The polypeptide and polynucleotide sequences have applications in
 CC diagnostics, forensics, gene mapping, identification of mutations
 CC responsible for genetic disorders or other traits to assess biodiversity
 CC and to produce other types of data and products dependent on DNA and
 CC amino acid sequences. AAS64197-AAS94564 represent novel human
 CC diagnostic coding sequences of the invention.
 CC Note: The sequence data for this patent did not appear in the printed
 CC specification, but was obtained in electronic format directly from WIPO
 CC at ftp.wipo.int/pub/published_pct_sequences.

XX Sequence 4977 BP: 1417 A: 1392 C: 1202 G: 966 T: 0 other:

Alignment Scores:
 Pred. No.: 795 Length: 4977
 Score: 41.00 Matches: 7
 Percent Similarity: 100.00% Conservative: 2
 Best Local Similarity: 77.78% Mismatches: 0
 Query Match: 93.16% Indels: 0
 DB: 23 Gaps: 0

US-09-965-594-1_COPY_14_22 (1-9) x AAS86668 (1-4977)

QY 1 LeuLeuGlyCysIleIleThrSerLeu 9

Db 1530 CTTTGGGCGTGTTCCTTACTTCCTG 1504

RESULT: 6

AAS88891/C

ID AAS88891 standard; cDNA: 4977 BP.

XX AAS88891;

XX 13-FEB-2002 (first entry)

XX DNA encoding novel human diagnostic protein #24595.

XX Human; chromosome mapping; gene mapping; gene therapy; forensic;
 KW food supplement; medical imaging; diagnostic; genetic disorder; ss.

XX Homo sapiens.

XX WO200175067-A2.

XX 11-OCT-2001.

XX 30-MAR-2001; 2001WO-US08631.

XX 31-MAR-2000; 2000US-0540217.

XX 23-AUG-2000; 2000US-0649167.

XX (HYSE-) HYSEQ INC.

XX Drmanac RT, Liu C, Tang YT;

XX WPI: 2001-639362/73.

DR P-PSDB; ABG24704.

XX New isolated polynucleotide and encoded polypeptides, useful in
 PT diagnostics, forensics, gene mapping, identification of mutations
 PT responsible for genetic disorders or other traits and to assess
 PT biodiversity.

XX Claim 1: SEQ ID No 24595; 103pp; English.

XX The invention relates to isolated polynucleotide (I) and

CC polypeptide (II) sequences. (II) is useful as hybridisation probes,

CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome

CC and gene mapping, and in recombinant production of (II). The
 CC polynucleotides are also used in diagnostics as expressed sequence tags
 CC for identifying expressed genes. (I) is useful in gene therapy techniques
 CC to restore normal activity of (II) or to treat disease states involving
 CC (II). (II) is useful for generating antibodies against it, detecting or
 CC quantitating a polypeptide in tissue, as molecular weight markers and as
 CC a food supplement. (II) and its binding partners are useful in medical
 CC imaging of sites expressing (II). (I) and (II) are useful for treating
 CC disorders involving aberrant protein expression or biological activity.
 CC the polypeptide and polynucleotide sequences have applications in
 CC diagnostics, forensics, gene mapping, identification of mutations
 CC responsible for genetic disorders or other traits to assess biodiversity
 CC and to produce other types of data and products dependent on DNA and
 CC amino acid sequences. AAS64197-AAS94564 represent novel human
 CC diagnostic coding sequences of the invention.
 CC Note: The sequence data for this patent did not appear in the printed
 CC specification, but was obtained in electronic format directly from WIPO
 CC at ftp.wipo.int/pub/published_pct_sequences.

XX Sequence 4977 BP: 1417 A: 1392 C: 1202 G: 966 T: 0 other:

Alignment Scores:
 Pred. No.: 795 Length: 4977
 Score: 41.00 Matches: 7
 Percent Similarity: 100.00% Conservative: 2
 Best Local Similarity: 77.78% Mismatches: 0
 Query Match: 93.16% Indels: 0
 DB: 23 Gaps: 0

US-09-965-594-1_COPY_14_22 (1-9) x AAS88891 (1-4977)

QY 1 LeuLeuGlyCysIleIleThrSerLeu 9

Db 1530 CTTTGGGCGTGTTCCTTACTTCCTG 1504

RESULT: 7

AAS89442/C

ID AAS89442 standard; cDNA: 4977 BP.

XX AAS89442;

XX 13-FEB-2002 (first entry)

XX DNA encoding novel human diagnostic protein #25646.

XX Human; chromosome mapping; gene mapping; gene therapy; forensic;
 KW food supplement; medical imaging; diagnostic; genetic disorder; ss.

XX Homo sapiens.

XX WO200175067-A2.

XX 11-OCT-2001.

XX 30-MAR-2001; 2001WO-US08631.

XX 31-MAR-2000; 2000US-0540217.

XX 23-AUG-2000; 2000US-0649167.

XX (HYSE-) HYSEQ INC.

XX Drmanac RT, Liu C, Tang YT;

XX WPI: 2001-639362/73.

DR P-PSDB; ABG25655.

XX New isolated polynucleotide and encoded polypeptides, useful in
 PT diagnostics, forensics, gene mapping, identification of mutations
 PT responsible for genetic disorders or other traits and to assess
 PT biodiversity.

XX Claim 1: SEQ ID No 25646; 103pp; English.

CC The invention relates to isolated polypeptide (I) and
 CC polypeptide (II) sequences. (I) is useful as hybridisation probes.
 CC polynucleotide sequences (III) primers, oligonucleotides, and for chromosome
 CC and gene mapping, and in recombinant production of (I). The
 CC polynucleotides are also used in diagnostic assays as expressed sequence tags
 CC for identifying expressed genes. (I) is useful in gene therapy techniques
 CC to restore normal activity of (II) or to treat disease states involving
 CC (II). (II) is useful for generating antibodies against its defects or
 CC quantitating a polypeptide in tissue, as molecular weight markers and as
 CC a food supplement. (I) and its binding partners are useful in medical
 CC imaging of sites expressing (II). (I) and (II) are useful for treating
 CC disorders involving aberrant protein expression or biological activity.
 CC The polypeptide and polynucleotide sequences have applications in
 CC diagnostics, forensics, gene mapping, identification of mutations
 CC responsible for genetic disorders or other traits to assess biodiversity
 CC and to produce other types of data and products dependent on DNA and
 CC amino acid sequences. AAS4137-AAS4564 represent novel human
 CC diagnostic coding sequences of the invention.
 CC Note: The sequence data for this patent did not appear in the printed
 CC specification, but was obtained in electronic format directly from W:PO
 CC at ftp.wipo.int/pub/published_pat_sequences.

XX Sequence 4977 BP; 1417 A; 1392 C; 1202 G; 966 T; 0 other;

Alignment Scores:
 Pred. No.: 795 Length: 4977
 Score: 41.00 Matches: 7
 Percent Similarity: 100.00% Conservative: 2
 Best Local Similarity: 77.78% Mismatches: 0
 Query Match: 93.18% Indels: 0
 DB: 23 Gaps: 0

US-09-965-594-1_copy_14_22 (1-9) x AAS89842 (1-4977)

Cy 1 LeuLeuGlyCysIleIleThrSerLeu 9
 |||||:|||||:|||||:|||||:
 Db 1530 CTTTTCGGGTGTTGTCACCTCTCTC 1504

RESULT 9

AAX24843
 ID AAX24843 standard; DNA: 9595 BP.

XX AAX24843;

XX 21-JUN-1999 (first entry)

DE Infectious hepatitis C virus genotype 1b strain HC-J4 genome.

XX HCV; infectious clone; infection; diagnosis; therapy; vaccine;
 KW screening; assay; antiviral; virucide; ss.

XX Hepatitis C virus.

FE Key Location/Qualifiers
 FT CDS 342..9374
 FT /*tag- a

PN W09904038-A2.

PD 28-JAN-1999.

XX 16-JUL-1998; 96W0-US14588.

PR 27-JAN-1996; 98JUS-C04416.

PR 18-JUL-1997; 97US-C053062.

XX (USSH) US DEPT HEALTH & HUMAN SERVICES.

XX Buxh J, Emerson SU, Purcell RH, Yanagi M;

XX WPI; 1999-192252/11.

DR P-PSDB; AAW95022.

XX

PT New isolated hepatitis C virus nucleic acids - used to develop
 PT products for the diagnosis, prevention and treatment of HCV
 PT infections and for developing screening assays

PS claim 3; Fig 1A-F; 126pp; English.

XX The present sequence comprises the nucleic acid sequence of the
 CC genome of infectious hepatitis C virus (HCV) genotype 1b strain
 CC HC-24 (ATCC 209596); that is capable of expressing this virus when
 CC transfected into cells. HC-J4 was obtained from acute phase plasma
 CC of a chimpanzee experimentally infected with serum containing
 CC HC-24/91. The claimed infectious nucleic acid sequence can be used
 CC to produce chimeric genomes (see AAX24843) consisting of the open
 CC reading frames of infectious nucleic acid sequences of other
 CC genotypes (including genotypes 1-6) and subtypes (such as 1b, 2a,
 CC 2b, 2c, 3a, 4a-f, 5a and 6a) of HCV. The invention also relates to
 CC the introduction of mutations or deletions into infectious nucleic
 CC acid sequences in order to produce an attenuated HCV virus suitable
 CC for vaccine development. Infectious nucleic acid sequences can
 CC also be used to produce attenuated virus via passage in vitro or in
 CC vivo of the viruses produced by transfection of a host cell with
 CC the infectious nucleic acid sequence. Vaccines comprising one or
 CC more polypeptides made from the infectious nucleic acid sequence are
 CC used to immunise mammals, especially humans, against hepatitis C.
 CC The nucleic acid sequences can also be used to induce protective
 CC immunity against the virus. The nucleic acid sequences or their
 CC encoded proteases (e.g. NS3 protease) can additionally be used to
 CC develop screening assays to identify antiviral agents for HCV.

XX Sequence 9595 BP; 1934 A; 2842 C; 2698 G; 2121 T; 0 other;

Alignment Scores:
 Pred. No.: 1-66e+03 Length: 9595
 Score: 41.00 Matches: 8
 Percent Similarity: 100.00% Conservative: 1
 Best Local Similarity: 88.89% Mismatches: 0
 Query Match: 93.18% Indels: 0
 DB: 20 Gaps: 0

US-09-965-594-1_COPY_14_22 (1-9) x AAX24843 (1-9595)

Cy 1 LeuLeuGlyCysIleIleThrSerLeu 9
 |||||:|||||:|||||:|||||:
 Db 3456 GTACITGGTTCATCATCACTAGCCCTC 3482

RESULT 9

AAC86939

ID AAC86939 standard; DNA: 9595 BP.

XX AAC86939;

XX 02-APR-2001 (first entry)

XX Nucleotide sequence of a hepatitis C virus (HCV) clone genotype 1b.

XX Chimeric virus; bovine viral diarrhoea virus; BVDV; hepatitis C virus;
 KW HCV; vaccine; viral inhibitor; antiviral; ss.

XX Hepatitis C virus.

FE Key Location/Qualifiers
 FT CDS 342..9374
 FT /*tag- a

PN W0200075352-A2.

XX 14-DEC-2000.

XX 02-JUN-2000; 2000W0-US15527.

XX 04-JUN-1999; 99US-0137917.

XX (USSH) US DEPT HEALTH & HUMAN SERVICES.

XX Nam J, Bukh J, Emerson SJ, Purcell RH;
 XX WPI: 2001-071061/C8.
 DR P-PSDB; AAB31170.

XX New nucleic acid comprising a chimeric bovine viral diarrhoea virus
 PT genome in which the (non-)structural region has been replaced by
 PT hepatitis C virus (HCV) genome useful for treating or preventing HCV
 PT signs and symptoms

XX Disclosure; Fig 4A-F; 97pp; English.

XX The specification describes a nucleic acid comprising a chimeric virus
 CC genome, specifically bovine viral diarrhoea virus (BVDV) genome in which
 CC the (non-)structural region has been replaced by the (non-)structural
 CC region of a hepatitis C virus (HCV) genome. The nucleic acids comprising
 CC the chimeric virus and the chimeric virus are useful for identifying
 CC cell lines capable of supporting the replication of these chimeric
 CC viruses, in screening for neutralizing antibodies to HCV of different
 CC genotypes, in the production of HCV-BVDV viruses, for the development
 CC of inactivated or attenuated vaccines to prevent HCV-BVDV in a mammal,
 CC in studying the molecular properties of HCV indirectly in vitro, and in
 CC identifying inhibitors of viral enzyme activity which would be useful
 CC as antiviral agents. Formulations of compositions comprising the
 CC chimeric viruses may be used to treat or prevent the signs and symptoms
 CC of HCV. The present sequence represents a BVDV clone, which is used
 CC to construct chimeric nucleic acids of the invention.

XX Sequence 9595 BP; 1934 A; 2843 C; 2697 G; 2121 T; 0 other;

Alignment Scores:
 Pred. No.: 1,660+03 Length: 9595
 Score: 41.00 Matches: 6
 Percent Similarity: 100.00% Conservatve: 1
 Best Local Similarity: 88.89% Mismatches: 0
 Query Match: 93.18% Indels: 0
 DB: 22 Gaps: 0

US-09-965-594-1_COPY_14_22 (1-9) x AAC66939 (1-9595)

Oy 1 LeuLeuGlyCysIleIleThrSerLeu 9

Db 3456 GTACTTGTTGCATCATCACTAGCCIC 3482

RESULT 10

AAF23492

ID AAF23492 standard; DNA; 9595 BP.

AC AAF23492;

XX 21-MAR-2001 (first entry)

DE Infectious Hepatitis C virus 1b genotype.

XX GBV-B; hepatitis C virus; HCV; vaccine; ds.

OS Hepatitis C virus.

XX WO200075337-A1.

PN 14-DEC-2000.

XX 02-JUN-2000; 2000WO-US15293.

XX 04-JUN-1999; 99US-0137694.

XX (USSS) US DEPT HEALTH & HUMAN SERVICES.

XX Bukh J, Yanagi M, Emerson SJ, Purcell RH;

XX WPI: 2001-091214/10.

XX

PT New infectious nucleic acids of the GB virus-B clone, useful for
 PT indirectly studying the molecular properties of hepatitis C virus (HCV)
 PT and in developing vaccines and therapeutics for HCV
 XX Disclosure; Fig 7; 96pp; English.

XX The present invention relates to GB virus-B. The nucleic acid molecules
 CC of the invention are useful for indirectly studying the molecular
 CC properties of hepatitis C virus (HCV). The infectious nucleic acid
 CC sequence of the GB virus-B clone and the HCV/GBV-B chimeras may be used
 CC in the development of vaccines and therapeutics for HCV.

XX Sequence 9595 BP; 1934 A; 2843 C; 2697 G; 2121 T; 0 other;

Alignment Scores:
 Pred. No.: 1,660+03 Length: 9595
 Score: 41.00 Matches: 6
 Percent Similarity: 100.00% Conservatve: 1
 Best Local Similarity: 88.89% Mismatches: 0
 Query Match: 93.18% Indels: 0
 DB: 22 Gaps: 0

US-09-965-594-1_COPY_14_22 (1-9) x AAF23492 (1-9595)

Oy 1 LeuLeuGlyCysIleIleThrSerLeu 9

Db 3456 GTACTTGTTGCATCATCACTAGCCIC 3482

RESULT 1:

AAF23492

ID AAF23492 standard; DNA; 94895 BP.

AC AAF23492;

XX 20-MAR-2001 (first entry)

XX BAC containing repeats from centromeres 1-4 #25.

XX Centromere; microsome; vector; ds.

OS Arabidopsis thaliana.

XX WO200055325-A2.

XX 21-SEP-2000.

XX 17-MAR-2000; 2000WO-US07392.

XX 16-MAR-1999; 99US-0125219.

XX 01-APR-1999; 99US-0127409.

XX 18-MAY-1999; 99US-0134770.

XX 13-SEP-1999; 99US-0153584.

XX 17-SEP-1999; 99US-0154603.

XX (UYCH) UNIV CHICAGO.

XX Preuss D, Copenhaver G, Keith K;

XX WPI: 2000-587529/55.

XX Recombinant DNA construct comprising a plant centromere, useful for
 PT producing stably inherited microsome which can serve as vectors for
 PT the construction of transgenic plant and animal cells -
 XX Claim 102; Page 804-819; 1449pp; English.

XX The present invention relates to a recombinant DNA construct of a plant
 CC (Arabidopsis thaliana) centromere. The constructs are useful for
 CC producing stably inherited microsome which can serve as vectors for
 CC the construction of transgenic plant and animal cells expressing
 CC selected proteins such as hormones, enzymes, interleukins, clotting
 CC factors, cytokines, antibodies, and growth factors.

XX

SQ Sequence 94895 BP: 28943 A; 18101 C; 18466 G; 29345 T; 0 other:

Alignment Scores:

Pred. No.:	2,2e+04	Length:	94895
Score:	41.00	Matches:	7
Percent Similarity:	100.00%	Conservative:	2
Best Local Similarity:	77.78%	Mismatches:	0
Query Match:	93.18%	Indels:	0
DB:	21	Gaps:	0

US-09-965-594-1_COPY_14_22 (1-9) x AAF22302 (1-94895)

Qy 1 LeuLeuGlyCysIleIleThrSerLeu 9
|||||:|||||:|||||:|||||:

Db 3572 TTCTTAGTTCGATTGTCGACGATG 3656

RESULT 12

AAF22302/C

ID AAF22302 standard; DNA: 94895 BP.

XX AC AAF22302;

XX DT 20-MAR-2001 (first entry)

XX DE BAC containing repeats from centromeres 1-4 #25.

XX KW Centromere; microsome; vector; ds.

XX OS Arabidopsis thaliana.

XX PN W0200055325-A2.

XX PD 21-SEP-2000.

XX PF 17-MAR-2000; 2006WO-US07392.

XX PR 18-MAR-1999; 99US-0125219.

XX PR 01-APR-1999; 99US-0127409.

XX PR 18-MAY-1999; 99US-0134770.

XX PR 13-SEP-1999; 99US-0153584.

XX PR 17-SEP-1999; 99US-0154603.

XX PA {CYCH-} UNIV CHICAGO.

XX PI Preuss D. Copenhaver G, Keith K;

XX DR WPI: 2000-587529/55.

XX PT Recombinant DNA construct comprising a plant centromere, useful for producing stably inherited microsome which can serve as vectors for the construction of transgenic plant and animal cells.

XX PS Claim 102; page 804-819; 1449pp; English.

XX CC The present invention relates to a recombinant DNA construct of a plant (Arabidopsis thaliana) centromere. The constructs are useful for producing stably inherited microsome which can serve as vectors for the construction of transgenic plant and animal cells expressing selected proteins such as hormones, enzymes, cytokines, clotting factors, cytokines, antibodies, and growth factors.

SQ Sequence 94895 BP: 28943 A; 18101 C; 18466 G; 29345 T; 0 other:

Alignment Scores:

Pred. No.:	2,2e+04	Length:	94895
Score:	41.00	Matches:	7
Percent Similarity:	100.00%	Conservative:	2
Best Local Similarity:	77.78%	Mismatches:	0
Query Match:	93.18%	Indels:	0
DB:	21	Gaps:	0

US-09-965-594-1_COPY_14_22 (1-9) x AAF22302 (1-94895)

Qy 1 LeuLeuGlyCysIleIleThrSerLeu 9
|||||:|||||:|||||:|||||:

Db 44375 TTCTTAGTTCGATTGTCGACGATG 44349

RESULT 13

AAF22287

ID AAF22287 standard; DNA: 103929 BP.

XX AC AAF22287;

XX DT 20-MAR-2001 (first entry)

XX DE BAC containing repeats from centromeres 1-4 #10.

XX KW Centromere; microsome; vector; ds.

XX OS Arabidopsis thaliana.

XX PN W0200055325-A2.

XX PD 21-SEP-2000.

XX PF 17-MAR-2000; 2006WO-US07392.

XX PR 18-MAR-1999; 99US-0125219.

XX PR 01-APR-1999; 99US-0127409.

XX PR 18-MAY-1999; 99US-0134770.

XX PR 13-SEP-1999; 99US-0153584.

XX PR 17-SEP-1999; 99US-0154603.

XX PA {CYCH-} UNIV CHICAGO.

XX PI Preuss D. Copenhaver G, Keith K;

XX DR WPI: 2000-587529/55.

XX PT Recombinant DNA construct comprising a plant centromere, useful for producing stably inherited microsome which can serve as vectors for the construction of transgenic plant and animal cells.

XX PS Claim 102; page 484-508; 1449pp; English.

XX CC The present invention relates to a recombinant DNA construct of a plant (Arabidopsis thaliana) centromere. The constructs are useful for producing stably inherited microsome which can serve as vectors for the construction of transgenic plant and animal cells expressing selected proteins such as hormones, enzymes, cytokines, clotting factors, cytokines, antibodies, and growth factors.

SQ Sequence 103929 BP: 32987 A; 19310 C; 18547 G; 34065 T; 0 other:

Alignment Scores:

Pred. No.:	2,43e+04	Length:	103929
Score:	41.00	Matches:	7
Percent Similarity:	100.00%	Conservative:	2
Best Local Similarity:	77.78%	Mismatches:	0
Query Match:	93.18%	Indels:	0
DB:	21	Gaps:	0

US-09-965-594-1_COPY_14_22 (1-9) x AAF22287 (1-103929)

Qy 1 LeuLeuGlyCysIleIleThrSerLeu 9
|||||:|||||:|||||:|||||:

Db 5120 TTCTTAGTTCGATTGTCGACGATG 5146

RESULT 14

AAF22286

ID AAF22286 standard; DNA: 134499 BP.

XX AC AAF22286;

XX DT 20-MAR-2001 (first entry)

DE BAC containing repeats from centromeres 1-4 #9.
 XX
 KW Centromere: microsome; vector; ds.
 XX
 XX Arabidopsis thaliana.
 OS
 PN WO200055325-A2.
 XX
 XX 21-SEP-2000.
 XX
 XX 17-MAR-2000; 2000WO-US07392.
 PF
 XX 18-MAR-1999; 99US-0125219.
 PR
 PR 01-APR-1999; 99US-0127409.
 PR
 PR 18-MAY-1999; 99US-0134770.
 PR
 PR 13-SEP-1999; 99US-0154584.
 PR
 PR 17-SEP-1999; 99US-0154603.
 XX
 XX (UYCH-) UNIV CHICAGO.
 PA
 XX Prouss D, Copenhaver G, Keith K;
 PI
 XX WPI: 2000-587529/55.
 DR
 XX Recombinant DNA construct comprising a plant centromere, useful for
 PT producing stably inherited microsome which can serve as vectors for
 PI the construction of transgenic plant and animal cells
 PT
 XX Claim 102; Page 451-484; 1449pp; English.
 PS
 XX The present invention relates to a recombinant DNA construct of a plant
 CC (Arabidopsis thaliana) centromere. The constructs are useful for
 CC producing stably inherited microsome which can serve as vectors for
 CC the construction of transgenic plant and animal cells expressing
 CC selected proteins such as hormones, enzymes, interleukins, clotting
 CC factors, cytokines, antibodies, and growth factors.
 CC
 XX Sequence 134499 BP; 41565 A; 25130 C; 25225 G; 42577 T; 2 other;
 SQ
 Alignment Scores:
 Pred. No.: 3-25e-04 Length: 134499
 Score: 41.00 Matches: 7
 Percent Similarity: 100.00% Conservatives: 2
 Best Local Similarity: 77.78% Mismatches: 0
 Query Match: 93.18% Indels: 0
 DB: 21 Gaps: 0
 US-09-965-594-1_COPY_14_22 (1-9) x AAF22286 (1-134499)
 QY 1 LeuleuGlyCysIleIleThrSerLeu 9
 DB 1360 TTGTTAGTTCATTGTCGACGAGATG 1386
 RESULT 15
 AAF22305
 ID AAF22305 standard; DNA: 1082138 BP.
 AC
 XX AAF22305;
 XX
 XX 20-MAR-2001 (first entry)
 DT
 XX Arabidopsis thaliana chromosome 4 centromere.
 DE
 XX Centromere: microsome; vector; ds.
 XX
 XX Arabidopsis thaliana.
 OS
 PN WO200055325-A2.
 XX
 XX 21-SEP-2000.
 XX
 XX 17-MAR-2000; 2000WO-US07392.
 PF
 XX 18-MAR-1999; 99US-0125219.
 PR
 PR 01-APR-1999; 99US-0127409.
 PR
 PR 18-MAY-1999; 99US-0134770.
 PR
 PR 13-SEP-1999; 99US-0154584.
 PR
 PR 17-SEP-1999; 99US-0154603.
 XX
 XX (UYCH-) UNIV CHICAGO.
 PA
 XX Prouss D, Copenhaver G, Keith K;
 PI
 XX WPI: 2000-587529/55.
 DR
 XX Recombinant DNA construct comprising a plant centromere, useful for
 PT producing stably inherited microsome which can serve as vectors for
 PI the construction of transgenic plant and animal cells
 PT
 XX Claim 102; Page 451-484; 1449pp; English.
 PS
 XX The present invention relates to a recombinant DNA construct of a plant
 CC (Arabidopsis thaliana) centromere. The constructs are useful for
 CC producing stably inherited microsome which can serve as vectors for
 CC the construction of transgenic plant and animal cells expressing
 CC selected proteins such as hormones, enzymes, interleukins, clotting
 CC factors, cytokines, antibodies, and growth factors.
 CC
 XX Sequence 134499 BP; 41565 A; 25130 C; 25225 G; 42577 T; 2 other;
 SQ

PR 18-MAR-1999; 99US-0125219.
 PR 01-APR-1999; 99US-0127409.
 PR 18-MAY-1999; 99US-0134770.
 PR 13-SEP-1999; 99US-0154584.
 PR 17-SEP-1999; 99US-0154603.
 XX
 XX (UYCH-) UNIV CHICAGO.
 PA
 XX Prouss D, Copenhaver G, Keith K;
 PI
 XX WPI: 2000-587529/55.
 DR
 XX Recombinant DNA construct comprising a plant centromere, useful for
 PT producing stably inherited microsome which can serve as vectors for
 PI the construction of transgenic plant and animal cells
 PT
 XX Claim 66; Page 977-1388; 1449pp; English.
 PS
 XX The present invention relates to a recombinant DNA construct of a plant
 CC (Arabidopsis thaliana) centromere. The constructs are useful for
 CC producing stably inherited microsome which can serve as vectors for
 CC the construction of transgenic plant and animal cells expressing
 CC selected proteins such as hormones, enzymes, interleukins, clotting
 CC factors, cytokines, antibodies, and growth factors.
 CC
 XX Sequence 1082138 BP; 348775 A; 194404 C; 195515 G; 343444 T; 0 other;
 SQ
 Alignment Scores:
 Pred. No.: 3-24e+05 Length: 1082138
 Score: 41.00 Matches: 7
 Percent Similarity: 100.00% Conservatives: 2
 Best Local Similarity: 77.78% Mismatches: 0
 Query Match: 93.18% Indels: 0
 DB: 21 Gaps: 0
 US-09-965-594-1_COPY_14_22 (1-9) x AAF22305 (1-1082138)
 QY 1 LeuleuGlyCysIleIleThrSerLeu 9
 DB 885359 TTGTAGCTTCATTGTCGACGAGATG 885065
 Search completed: September 29, 2003, 19:19:33
 Job time : 244 secs

GenCore version 5.1.1
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM protein - nucleic search, using frame_plus_p2a_role1

Run on: September 29, 2003, 19:11:08 : Search time 1625 seconds
(without alignments)
134,539 William Cole updates/sec

Title: US-09-965-594-1_COPY_14_22

Perfect score: 44

Sequence: 1 LACIITS:9

Scoring table: BLOSUM62

Xgapop 10.0 , Xgapext 0.5

Ygapop 10.0 , Ygapext 0.5

Fgapop 6.0 , Fgapext 7.0

Delop 5.0 , Delext 7.0

Searched: 22781392 seqs, 12152238056 residues

Total number of hits satisfying chosen parameters: 45562778

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 99%

Listing first 45 summaries

Command line parameters:

-MODEL=frame+P2n.model -DEV=xit
-Q=/cgn2_1/USFOO-pool/US09965594/runat_29092003_164249_14082/app_query.fasta_1.59
-DB=EST -QFMT=fastap -SUFFIX=rst -MINMATCH=0.1 -LOOPCL=0 -LOOPEXT=0
-UNITS=bits -SPART=1 -MATRIX=blosum62 -TRANS=human40.cdi -LIST=45
-DOCALIGN=200 -THR_SCORE=pct -THR_MAX=99.9 -THR_MIN=0 -ALIGN=15 -MODE=LOCAL
-OUTFMT=ptc -NORM=ext -HEAPSIZE=500 -MINLEN=0 -MAXLEN=200000000
-USER=US09965594 -ACGN_1_1_2810 -runat_29092003_164249_14082 -NUPU=6 -ICPU=3
-NO_MMAP -LARGEQUERY -NEG_SCORES=0 -WAIT -DSPBLOCK=100 -LONGLOG
-DEV_TIMEOUT=120 -WARN_TIMEOUT=30 -THREADS=1 -XGAPOP=10 -XGAPEXT=0.5 -FGAPOP=6
-FGAPEXT=7 -YGAPOPOP=10 -YGAPEXT=0.5 -DELOP=6 -DELEXT=7

Database :

- 1: em_estab:*
- 2: cm_esthum:*
- 3: cm_estab:*
- 4: cm_estab:*
- 5: em_estab:*
- 6: em_estab:*
- 7: em_estab:*
- 8: cm_hic:*
- 9: qb_est1:*
- 10: qb_est2:*
- 11: qb_hic:*
- 12: qb_est3:*
- 13: qb_est4:*
- 14: qb_est5:*
- 15: cm_esthum:*
- 16: em_estom:*
- 17: cm_gss_hum:*
- 18: cm_gss_inv:*
- 19: em_gss_plc:*
- 20: em_gss_vrt:*
- 21: em_gss_fuc:*
- 22: em_gss_mam:*
- 23: em_gss_mus:*
- 24: cm_gss_pro:*
- 25: em_gss_rod:*
- 26: em_gss_phg:*
- 27: em_gss_vrl:*
- 28: qb_gss1:*

29: gb_gss2:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	% Match	Query Length	DB ID	Description
C 1	41	93.2	168	BH241427	BH241427 AUILF55TR
C 2	41	93.2	240	AL950918	AL950918 A-abidops
C 3	41	93.2	334	BZ352875	BZ352875 SALK_0827
C 4	41	93.2	400	CNS00021	AL085272 Arabidops
C 5	41	93.2	424	BH758372	BH758372 SALK_0185
C 6	41	93.2	444	BF651538	BF651538 SALK_0185
C 7	41	93.2	465	BF651537	BF651537 274341 MA
C 8	41	93.2	498	B96162	B96162 F26H1TFB IG
C 9	41	93.2	505	B26036	B26036 J532TF TAWU
C 10	41	93.2	631	BZ289143	BZ289143 SALK_0225
C 11	41	93.2	683	BH241556	BH241556 AUILF55TR
C 12	41	93.2	693	B95974	B95974 F21D4TFC IG
C 13	41	93.2	719	BH241494	BH241494 AUILF22TR
C 14	41	93.2	724	AO955329	AO955329 LERAD34TF
C 15	41	93.2	758	BH241165	BH241165 AUILF85TF
C 16	41	93.2	787	BH241117	BH241117 AUILF85TF
C 17	41	93.2	823	BH241591	BH241591 AUILF55TF
C 18	41	93.2	831	BH241228	BH241228 AUILF15TF
C 19	41	93.2	835	CC429018	CC429018 PUHJV59TD
C 20	41	93.2	847	CC363288	CC363288 PUHJ110TD
C 21	41	93.2	917	BF167496	BF167496 601775095
C 22	40	90.9	287	AI800243	AI800243 t176d09.x
C 23	40	90.9	309	AI587235	AI587235 tq21f04.x
C 24	40	90.9	355	14	Z39665 HSC1GA052 n
C 25	40	90.9	384	9	AI583735 tt03h01.x
C 26	40	90.9	401	14	CA902724 PCS01861X
C 27	40	90.9	406	10	BF109646 7166c06.x
C 28	40	90.9	407	10	BF727441 ky21s011.x
C 29	40	90.9	413	28	AQ759141 HS_2201.A
C 30	40	90.9	423	9	AA721993 zh17d10.s
C 31	40	90.9	424	14	TC3709 IB779 infan
C 32	40	90.9	425	9	AA812004 OB39e08.s
C 33	40	90.9	426	9	AW665172 h187q11.x
C 34	40	90.9	473	9	AI927322 w089e11.x
C 35	40	90.9	540	12	BQ017316 ul-h-DTI-
C 36	40	90.9	588	10	BE785099 601478355
C 37	40	90.9	588	14	H15452 ym29g06.s1
C 38	40	90.9	613	14	H15273 ym30e09.s1
C 39	40	90.9	625	12	BM682281 UI-E-EJ0-
C 40	40	90.9	683	29	BZ325209 IC50e11.g
C 41	40	90.9	702	13	BQ184273 UI-E-EJ1-
C 42	40	90.9	710	13	BQ187309 UI-E-EJ1-
C 43	40	90.9	915	10	BG685367 602637211
C 44	40	90.9	927	13	BU527696 AGENCOURT
C 45	40	90.9	1006	10	BG758345 602712536

ALIGNMENTS

RESULT 1
BH241427/c
LOCUS
DEFINITION BH241427 AUILF55TR AUIL Arabidopsis thaliana genomic clone AUILF55, genomic survey sequence.
ACCESSION BH241427
VERSION BH241427.1 GI:16915855
KEYWORDS GSS
SOURCE Arabidopsis thaliana (thale cress)
ORGANISM Arabidopsis thaliana
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids
; eurosids II; Brassicales; Brassicaceae; Arabidopsi.

```

REFERENCE
AUTHORS      1 (bases 1 to 106);
              Feldblyum,T.V. and Fraser,C.M.
TITLE        Survey sequencing of Arabidopsis thaliana BAC F26J21
JOURNAL      Unpublished
COMMENT      Other_GSSs: AU1LF55TF
              Contact: Chris Town
              TIGR
              3712 Medical Center Drive, Rockville, MD 20850, USA.
              Tel: 301-838-3523
              Fax: 301-838-0208
              Email: cdtown@tigr.org
              Prom Wash. U contig 720.
              Seq primer: 1R
              Class: sheared ends.
FEATURES
source       Location/Qualifiers
              1..108
              /organism="Arabidopsis thaliana"
              /mol_type="genomic DNA"
              /strain="Columbia"
              /db_xref="taxon:3702"
              /clone="AU1LF55"
              /clone_lib="AUIU"
              /note="Vector: pHC82; Site 1: BstXI; 2-3 kb sheared BAC
              DNA inserted into pHC82 using BstXI linkers"
BASE COUNT   36 a    29 c    8 g    35 t
ORIGIN
Alignment Scores:
Pred. No.:      90.9      Length:      106
Score:          41.00     Matches:      7
Percent Similarity: 100.00% Conservative: 2
Best Local Similarity: 77.78% Mismatches: 0
Query Match:     93.18% Indels:      0
DB:             28      Gaps:       0

US-09-965-594-1_COPY_14_22 (1-9) x B241427 (1-103)

QY 1 LeuLeuGlyCysIleIleThrSerLeu 9
   |||||
DB 56 TTGTTAGGTGCATTCGTGACGAGTATG 39

RESULT 2
AL950918      230 bp      DNA      linear      GSS 24-OCT-2002
LOCUS        Arabidopsis thaliana T-DNA flanking sequence GK-332C10-016064,
DEFINITION   genomic survey sequence.
ACCESSION    AL950918.1 GI:24407540
VERSION      AL950918.1
KEYWORDS     GSS.
SOURCE       Arabidopsis thaliana (thale cress)
ORGANISM     Arabidopsis thaliana
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
rosids; eutosids II; Brassicales; Brassicaceae; Arabidopsis.
REFERENCE    1
AUTHORS      Strizhov,N., Li,Y., Rosso,M., Viehoever,P., Dekker,K., Saedler,H.
              and Weissshaar,B.
TITLE        A pipeline for automated high-throughput generation of FSTs
              (flanking sequence tags) from Arabidopsis thaliana T-DNA
              transformed lines
JOURNAL      Unpublished
REFERENCE    2
AUTHORS      Rosso,M., Strizhov,N., Li,Y., Reiss,B., Dekker,K. and Weissshaar,B.
TITLE        A new Arabidopsis thaliana T-DNA mutagenised population (GABI-Kat)
              for flanking sequence tag based reverse genetics
JOURNAL      Unpublished
REFERENCE    3 (bases 1 to 230)
AUTHORS      Li,Y., Rosso,M., Strizhov,N. and Weissshaar,B.
TITLE        Direct Submission
JOURNAL      Submitted (21-OCT-2002) Weissshaar B., Max-Planck-Institut fuer
              Zuechtungsforschung, Carl-von-Linne-Weg 10, Koeln, 50829, Germany
COMMENT      This sequence is recovered from the left border of the T-DNA. It

```

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indicates an insertion within the locus defined by clone F4M19. The
sequences are generated at the MPI for Plant Breeding Research in
the context of the GABI-Kat project. GABI-Kat is part of the German
Plant Genomics program designated "GABI". Information on line
availability can be found at:
http://www.mpiz-koeln.mpg.de/GABI-Kat/.
FEATURES
source       Location/Qualifiers
              1..230
              /organism="Arabidopsis thaliana"
              /mol_type="genomic DNA"
              /strain="Columbia 0"
              /db_xref="taxon:3702"
              /clone="GK-332C10-016064"
              /clone_lib="Arabidopsis thaliana T-DNA insertion lines"
              /note="PCR was performed on DNA from Arabidopsis thaliana
              plants (T1) which were transformed with the T-DNA from
              vector PAC161. The lines contain one or more T-DNA
              insertions. The DNA fragment(s) resulting from the PCR
              were directly sequenced to determine the genomic sequence
              flanking the insertion. Sequences displaying significant
              similarity to the A. thaliana nuclear genome sequence were
              processed for submission. T-DNA derived sequences were
              removed"
BASE COUNT   67 a    42 c    48 g    73 t
ORIGIN
Alignment Scores:
Pred. No.:      277      Length:      230
Score:          41.00     Matches:      7
Percent Similarity: 100.00% Conservative: 2
Best Local Similarity: 77.78% Mismatches: 0
Query Match:     93.18% Indels:      0
DB:             29      Gaps:       0

US-09-965-594-1_COPY_14_22 (1-9) x AL950918 (1-230)

QY 1 LeuLeuGlyCysIleIleThrSerLeu 9
   |||||
DB 104 TTGTTAGGTGCATTCGTGACGAGTATG 190

RESULT 3
B2352875/c    334 bp      DNA      linear      GSS 14-NOV-2002
LOCUS        SALK_082734.11.45.x Arabidopsis thaliana T-DNA insertion lines
DEFINITION   Arabidopsis thaliana genomic clone SALK_082734.31.45.x, genomic
              survey sequence.
ACCESSION    B2352875
VERSION      B2352875.1 GI:24943737
KEYWORDS     GSS.
SOURCE       Arabidopsis thaliana (thale cress)
ORGANISM     Arabidopsis thaliana
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids
; eucosids II; Brassicales; Brassicaceae; Arabidopsis.
REFERENCE    1 (bases 1 to 334)
AUTHORS      Alonso,J.M., Leisse,T.J., Harejas,P., Chen,H., Cheuk,R., Gadrinab
              ,C., Jeske,A., Karnes,M., Kim,C.J., Parker,H., Prednis,L., Shinn,P.
              , Zimmerman,J. and Ecker,J.K.
TITLE        A Sequence-Indexed Library of Insertion Mutations in the
              Arabidopsis Genome
JOURNAL      Unpublished
COMMENT      Contact: Joseph R. Ecker
              Salk Institute Genomic Analysis Laboratory (SIGNAL)
              The Salk Institute for Biological Studies
              10010 N. Torrey Pines Road, La Jolla, CA 92037, USA
              Tel: 858 453 4100 x1752
              Fax: 858 558 6379
              Email: ecker@salk.edu
              This is single pass sequence recovered from the left border of
              T-DNA.
              Class: T-DNA tagged.
              Location/Qualifiers
              1..334
FEATURES
source

```

```

/organism="Arabidopsis thaliana"
/mol_type="genomic DNA"
/strain="Columbia 0"
/db_xref="taxon:3702"
/clone="SALK_082734.1-1.45.x"
/notes="PCR was performed on Arabidopsis thaliana lines
each of which contains one or more TDNA insertion
elements. The resultant fragment for each line was
directly sequenced to determine the genomic sequence at
the site of insertion. Details of the protocols used can
be found at: http://signal.salk.edu/dna_protocols.html"
BASE COUNT 111 a 71 c 60 g 51 t
ORIGIN
Alignment Scores:
Pred. No.: 480 Length: 334
Score: 41.00 Matches: 7
Percent Similarity: 100.00% Conservative: 2
Best Local Similarity: 77.78% Mismatches: 0
Query Match: 93.18% Indels: 0
Gaps: 0
DB: 29

US-09-965-594-1_COPY_14_22 (1-9) x B2352875 (1-334)
Qy 1 LeuLeuGlyCysIleIleThrSerLeu 9
|||||:|||||:|||||:|||||:
Db 188 TTGTTAGGTGCATTGTGCAGGATG 162

RESULT 4
CNS00021/c
LOCUS
DEFINITION
Arabidopsis thaliana genome survey sequence T7 end of BAC F10H2 of
IGF library from strain Columbia of Arabidopsis thaliana, genomic
survey sequence.
ACCESSION
Al085272
VERSION
Al085272.1 GI:5286412
KEYWORDS
GSS.
SOURCE
Arabidopsis thaliana (thale cress)
ORGANISM
Arabidopsis thaliana
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
rosids; eurosids I; Brassicales; Brassicaceae; Arabidopsi.
1 (bases 1 to 400)
Sa.ancubati; M. Chaisne; N. Artiguenave; F. Brottier; P. Wincker; P.
Samson; D. Saurin; W. Weissbach; J. and Quetier; F.
Unpublished
Genoscope.
Direct Submission
Submitted (25-JUN-1999) Genoscope - Centre National de Sequence
BP 191 91006 Evry cedex - FRANCE (E-mail: seq@genoscope.cns.fr)
Web : www.genoscope.cns.fr
FEATURES
Location/Qualifiers
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/mol_type="genomic DNA"
/strain="Columbia"
/db_xref="taxon:3702"
/clone="F10H2"
/notes="end : T7"
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Alignment Scores:
Pred. No.: 627 Length: 400
Score: 41.00 Matches: 7
Percent Similarity: 100.00% Conservative: 2
Best Local Similarity: 77.78% Mismatches: 0
Query Match: 93.18% Indels: 0
Gaps: 0
DB: 29

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US-09-965-594-1_COPY_14_22 (1-9) x CNS00021 (1-400)
Qy 1 LeuLeuGlyCysIleIleThrSerLeu 9
|||||:|||||:|||||:|||||:
Db 350 TTGTTAGGTGCATTGTGCAGGATG 324

RESULT 5
BH758372
LOCUS
DEFINITION
SALK_018581.33.10.x Arabidopsis thaliana TDNA insertion lines
Arabidopsis thaliana genomic clone SALK_018581.33.10.x, genomic
survey sequence.
ACCESSION
BH758372
VERSION
BH758372.1 GI:19043622
KEYWORDS
GSS.
SOURCE
Arabidopsis thaliana (thale cress)
ORGANISM
Arabidopsis thaliana
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids;
eurosids II; Brassicales; Brassicaceae; Arabidopsi.
1 (bases 1 to 424)
Alonso; J.M.; Leisse; T.J.; Barajas; P.; Chen; H.; Cheuk; R.; Gadrinab;
C.; Jeske; A.; Karnes; M.; Kim; C.J.; Parker; H.; Prednis; L.; Shinn; P.;
Zimmerman; J. and Ecker; J.R.
A sequence-indexed library of insertion mutations in the
Arabidopsis Genome
Unpublished
Contact: Joseph R. Ecker
Salk Institute Genomic Analysis Laboratory (SIGAL)
The Salk Institute for Biological Studies
10010 N. Torrey Pines Road, La Jolla, CA 92037, USA
Tel: 858 453 4100 x1752
Fax: 858 558 6379
Email: ecker@salk.edu
This is single pass sequence recovered from the left border of
TDNA.
Class: TDNA tagged.
FEATURES
Location/Qualifiers
source
1..424
/organism="Arabidopsis thaliana"
/mol_type="genomic DNA"
/strain="Columbia 0"
/db_xref="taxon:3702"
/clone="SALK_018581.33.10.x"
/notes="PCR was performed on Arabidopsis thaliana lines
each of which contains one or more TDNA insertion
elements. The resultant fragment for each line was
directly sequenced to determine the genomic sequence at
the site of insertion. Details of the protocols used can
be found at: http://signal.salk.edu/dna_protocols.html"
BASE COUNT 62 a 151 c 98 g 113 t
ORIGIN
Alignment Scores:
Pred. No.: 683 Length: 424
Score: 41.00 Matches: 7
Percent Similarity: 100.00% Conservative: 2
Best Local Similarity: 77.78% Mismatches: 0
Query Match: 93.18% Indels: 0
Gaps: 0
DB: 28

US-09-965-594-1_COPY_14_22 (1-9) x BH758372 (1-424)
Qy 1 LeuLeuGlyCysIleIleThrSerLeu 9
|||||:|||||:|||||:|||||:
Db 74 TTGTTAGGTGCATTGTGCAGGATG 100

RESULT 6
BF651538/c
LOCUS
DEFINITION
274341 MARC 3B0V Bos taurus cDNA 5', rRNA sequence.
ACCESSION
BF651538

```

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VERSION      BF651538.1  GI:11915668
KEYWORDS
SOURCE       Bos taurus (cow)
ORGANISM     Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
             Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
             Bovidae; Bovinae; Bos.

REFERENCE    1 (bases 1 to 444)
AUTHORS      Smith,T.P.L., Grosse,W.M., Freking,B.A., Roberts,A.J., Stone,R.T.,
             Casas,E., Wray,J.E., White,J., Cho,J., Fahrenkrug,S.C., Bennett
             ,G.L., Heaton,M.P., Laegreid,W.W., Rohrer,G.A., Chitko-McKown,C.G.,
             Pertea,G., Holt,I., Karamycheva,S., Liang,F., Quackenbush,J. and
             Keele,J.W.
             Sequence evaluation of four pooled-tissue normalized bovine cDNA
             libraries and construction of a gene index for cattle
             Genome Res. 11 (4), 626-630 (2001)
             21180013
             PUBMED 11282378
             COMMENT Contact: Smith TPL
             USDA, ARS, US Meat Animal Research Center
             PO Box 166, Clay Center, NE 68933-0166, USA
             Tel: 402 762 4366
             Fax: 402 762 4396
             Email: smith@mail.marc.usda.gov
             Single pass sequencing. Bases called and alt_trimmed with phred
             v0.980904.c. Vector identified by cross_match with the -minscore 18
             and -mismatch 12 options.
             PCR Primers
             FORWARD: AGGAACAGCTATGACCAT
             BACKWARD: GTTTCCTCAGTCAGCAGC
             Plate: 53 row: B column: 22
             Seq primer: ATTAGTGACACTATAG.
             Location/Qualifiers
             source          :
             .. 444
             /organism="Bos taurus"
             /mol_type="mRNA"
             /db_xref="taxon:9913"
             /tissue_type="pooled"
             /lab_host="DH10B"
             /clone_lib="MARC 3BOV"
             /note="Vector: pCMV SPORT6; Site.1: NotI; Site.2: SalI;
             library made from pooled tissue from marrow, alveolar
             macrophage, ovary, fetal semitendinosus muscle, and fetal
             longissimus muscle."
             .. 444

BASE COUNT   116 a 96 c 115 g 117 t
ORIGIN
Alignment Scores:
Pred. No.: 731 Length: 444
Score: 41.00 Matches: 8
Percent Similarity: 100.00% Conservative: 1
Best Local Similarity: 88.89% Mismatches: 0
Query Match: 93.18% Indels: 0
DB: 10 Gaps: 0

US-09-965-594-l_COPY_14_22 (1-9) x BF651538 (1-444)
QY 1 LeuLeuGlyCysIleIleThrSerLeu 9
||||:|||||
DB 113 TTGGTGGTTCGTATAATTAATTAATTAATTA 87

RESULT 7
BF651537/c
LOCUS       BF651537
DEFINITION  274340 MARC 3BOV Bos taurus cDNA 5', mRNA sequence.
ACCESSION   BF651537
VERSION     BF651537.1 GI:11915667
KEYWORDS
SOURCE      Bos taurus (cow)
ORGANISM    Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
             Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
             Bovidae; Bovinae; Bos.

```

```

REFERENCE    1 (bases 1 to 465)
AUTHORS      Smith,T.P.L., Grosse,W.M., Freking,B.A., Roberts,A.J., Stone,R.T.,
             Casas,E., Wray,J.E., White,J., Cho,J., Fahrenkrug,S.C., Bennett
             ,G.L., Heaton,M.P., Laegreid,W.W., Rohrer,G.A., Chitko-McKown,C.G.,
             Pertea,G., Holt,I., Karamycheva,S., Liang,F., Quackenbush,J. and
             Keele,J.W.
             Sequence evaluation of four pooled-tissue normalized bovine cDNA
             libraries and construction of a gene index for cattle
             Genome Res. 11 (4), 626-630 (2001)
             21180013
             PUBMED 11282378
             COMMENT Contact: Smith TPL
             USDA, ARS, US Meat Animal Research Center
             PO Box 166, Clay Center, NE 68933-0166, USA
             Tel: 402 762 4366
             Fax: 402 762 4396
             Email: smith@mail.marc.usda.gov
             Single pass sequencing. Bases called and alt_trimmed with phred
             v0.980904.c. Vector identified by cross_match with the -minscore 18
             and -mismatch 12 options.
             PCR Primers
             FORWARD: AGGAACAGCTATGACCAT
             BACKWARD: GTTTCCTCAGTCAGCAGC
             Plate: 53 row: B column: 21
             Seq primer: ATTAGTGACACTATAG.
             Location/Qualifiers
             source          :
             .. 465
             /organism="Bos taurus"
             /mol_type="mRNA"
             /db_xref="taxon:9913"
             /tissue_type="pooled"
             /lab_host="DH10B"
             /clone_lib="MARC 3BOV"
             /note="Vector: pCMV SPORT6; Site.1: NotI; Site.2: SalI;
             library made from pooled tissue from marrow, alveolar
             macrophage, ovary, fetal semitendinosus muscle, and fetal
             longissimus muscle."
             .. 465

BASE COUNT   121 a 97 c 122 g 125 t
ORIGIN
Alignment Scores:
Pred. No.: 782 Length: 465
Score: 41.00 Matches: 8
Percent Similarity: 100.00% Conservative: 1
Best Local Similarity: 88.89% Mismatches: 0
Query Match: 93.18% Indels: 0
DB: 10 Gaps: 0

US-09-965-594-l_COPY_14_22 (1-9) x BF651537 (1-465)
QY 1 LeuLeuGlyCysIleIleThrSerLeu 9
||||:|||||
DB 113 TTGGTGGTTCGTATAATTAATTAATTAATTA 87

RESULT 8
BF6162/c
LOCUS       BF6162
DEFINITION  F26H1TFB IGF Arabidopsis thaliana genomic clone F26H1, genomic
             survey sequence.
ACCESSION   B96162
VERSION     B96162.1 GI:29986638
KEYWORDS
SOURCE      Arabidopsis thaliana (thale cress)
ORGANISM     Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
             Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids
             ; eurosids II; Brassicales; Brassicaceae; Arabidopsi.
             1 (bases 1 to 498)
             Rounsley,S.D., Field,C.E., Bass,S., Linher,K., Linher,K., Golden,K.
             , Berry,K., Granger,D., Suh,E., Wible,C., Adams,M.D. and Venter
             ,J.C.
             A BAC End Sequence Database for Identifying Minimal Overlaps in
             Arabidopsis Genomic Sequencing. Update 3

```



```

JOURNAL      Unpublished
COMMENT      Other_GSSs: F26H1R
             Contact: Steve Rounsley
             Department of Eukaryotic Genomics
             The Institute for Genomic Research
             9712 Medical Center Dr., Rockville, MD 20850, USA
             Tel: 301 838 0200
             Fax: 301 838 0208
             Email: rounsley@tigr.org
             Seq primer: M13-21
             Class: SAC ends
             High quality sequence stop: 498.
FEATURES     Location/Qualifiers
             ..498
             /organism="Arabidopsis thaliana"
             /mol_type="genomic DNA"
             /strain="Columbia"
             /db_xref="taxon:3702"
             /clone="F26H1"
             /sex="hermaphrodite"
             /note="Vector: pClOACII; Site_1: EcoRI; Site_2: EcoRI;
             Produced by Thomas Altmann"
BASE COUNT   184 a 105 c 80 g 129 t
ORIGIN

Alignment Scores:
Pred. No.:      866      Length:      498
Score:          41.00    Matches:      7
Percent Similarity: 100.00%  Conservative: 2
Best Local Similarity: 77.78%  Mismatches: 0
Query Match:      93.18%  Indels: 0
DB:               28      Gaps: 0

US-09-965-594-1_COPY_14_22 (1-9) x B96162 (1-498)

Qy 1 LeuLeuGlyCysIleIleThrSerLeu 9
|||||.....|.....|.....|.....|
Db 370 TTGTTAGGTTGCATTGTGACGAGTATG 344

RESULT 9
B26036/c  B26036  505 bp  DNA  linear  GSS 13-OCT-1997
LOCUS     B26036  505 bp  DNA  linear  GSS 13-OCT-1997
DEFINITION Arabidopsis thaliana genomic clone F5G2, genomic survey
sequence.
ACCESSION B26036
VERSION   B26036.1  GI:2512002
KEYWORDS  Arabidopsis thaliana (thale cress)
SOURCE    Arabidopsis thaliana
ORGANISM  Arabidopsis thaliana
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids
; eurosids II; Brassicales; Brassicaceae; Arabidopsis.
1 (bases 1 to 505)
Alonso,J.M., Leisse,T.J., Barajas,P., Chen,H., Cheuk,R., Gadrinab
,C., Jeske,A., Karnes,M., Kim,C.J., Parker,H., Predslis,L., Shinn,P.
,Zimmerman,J. and Ecker,J.R.
A Sequence-Indexed Library of Insertion Mutations in the
Arabidopsis Genome
Unpublished
Contact: Joseph R. Ecker
Salk Institute Genomic Analysis Laboratory (SIGNAL)
The Salk Institute for Biological Studies
10010 N. Torrey Pines Road, La Jolla, CA 92037, USA
Tel: 858 453 4100 x1752
Fax: 858 558 6379
Email: ecker@salk.edu
This is single pass sequence recovered from the left border of
TDNA.
Class: TDNA tagged.
Location/Qualifiers
..655
/organism="Arabidopsis thaliana"
/mol_type="genomic DNA"
/strain="Columbia 0"
/db_xref="taxon:3702"
/clone="SALK_022540.51-60.x"
/clone_lib="Arabidopsis thaliana TDNA insertion lines"
/note="PCR was performed on Arabidopsis thaliana lines
each of which contains one or more TDNA insertion
elements. The resultant fragment for each line was
directly sequenced to determine the genomic sequence at
the site of insertion. Details of the protocols used can
be found at http://signal.salk.edu/tdna_protocols.html"
BASE COUNT   207 a 131 c 113 g 197 t
ORIGIN

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Alignment Scores:
Pred. No.: 1.29e+03 Length: 651
Score: 41.00 Matches: 7
Percent Similarity: 100.00% Conservative: 2
Best Local Similarity: 77.78% Mismatches: 0
Query Match: 93.18% Indels: 0
DB: 29 Gaps: 0

US-09-965-594-1_COPY_14_22 (1-9) x B2289143 (1-651)
QY 1 LeuLeuGlyCysIleIleThrSerLeu 9
|||||
Db 170 TTGTAGTTCATTCGACGAGTATG 144

RESULT 11
BH241556 683 bp DNA linear GSS 13-NOV-2001
LOCUS AU1LF22TF AUIL Arabidopsis thaliana genomic clone AU1LF22, genomic
DEFINITION survey sequence.
ACCESSION BH241556
VERSION BH241556 GI:16916061
KEYWORDS Arabidopsis thaliana (thale cress)
SOURCE Arabidopsis thaliana
ORGANISM Arabidopsis thaliana
Eukaryota: Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids
; eucosids II; Brassicales; Brassicaceae; Arabidopsi.
REFERENCE 1 (bases 1 to 683)
AUTHORS Town,C.D., Whitelaw,C.A., Pai,G., Van Aken,S.E., Utterback,T.V.,
Feldblyum,T.V. and Fraser,C.M.
TITLE Survey sequencing of Arabidopsis thaliana BAC F26J2;
JOURNAL Unpublished
COMMENT Other_GSSs: AU1LF27TR
Contact: Chris Town
TIGR
9712 Medical Center Drive, Rockville, MD 20850, USA.
Tel: 301-838-3521
Fax: 301-838-0208
Email: cdtown@tigr.org
From Wash. U contig 720.
Seq primer: TF
Class: sheared ends.
Location/Qualifiers
source
1..683
/organism="Arabidopsis thaliana"
/mol_type="genomic DNA"
/strain="Columbia"
/db_xref="taxon:3702"
/clone="AU1577"
/clone_lib="IGF"
/notes="Vector: pPOS2; Site_1: BstXI; 2-3 kb sheared BAC
DNA inserted into pPOS2 using BstXI linkers"

BASE COUNT 196 a 129 c 138 g 220 t
ORIGIN

Alignment Scores:
Pred. No.: 1.38e+03 Length: 683
Score: 41.00 Matches: 7
Percent Similarity: 100.00% Conservative: 2
Best Local Similarity: 77.78% Mismatches: 0
Query Match: 93.18% Indels: 0
DB: 28 Gaps: 0

US-09-965-594-1_COPY_14_22 (1-9) x BH241556 (1-683)
QY 1 LeuLeuGlyCysIleIleThrSerLeu 9
|||||
Db 655 TTGTAGTTCATTCGACGAGTATG 681

RESULT 12
BH241556 693 bp DNA linear GSS 31-MAR-1998
LOCUS AU1LF22TF AUIL Arabidopsis thaliana genomic clone F21D4, genomic
DEFINITION survey sequence.
ACCESSION BH241556
VERSION BH241556 GI:16915962
KEYWORDS Arabidopsis thaliana (thale cress)
SOURCE Arabidopsis thaliana
ORGANISM Arabidopsis thaliana
Eukaryota: Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids
; eucosids II; Brassicales; Brassicaceae; Arabidopsi.
REFERENCE 1 (bases 1 to 693)
AUTHORS Town,C.D., Whitelaw,C.A., Pai,G., Van Aken,S.E., Utterback,T.V.,
Feldblyum,T.V. and Fraser,C.M.
TITLE Survey sequencing of Arabidopsis thaliana BAC F26J2;
JOURNAL Unpublished
COMMENT Other_GSSs: AU1LF22TF

```

Contact: Chris Town
TIGR
9712 Medical Center Drive, Rockville, MD 20850, USA.
Tel: 301-838-3523
Fax: 301-838-0208
Email: cdtown@tigr.org
From Wash. U contig 720.
Seq primer: TF
Class: Sheared ends.

FEATURES

source
Location/Qualifiers

1..719
/organism="Arabidopsis thaliana"
/mol_type="genomic DNA"
/strain="Columbia"
/db_xref="taxon:3702"
/clone_lib="AULF85"
/clone="AULF85"
/note="Vector: pPOS2; Site 1: BstXI; 2-3 kb sheared BAC DNA inserted into pPOS2 using BstXI linkers"

BASE COUNT 187 a 149 c 140 g 233 t

ORIGIN

Alignment Scores:
Pred. No.: 1,49e-03 Length: 719
Score: 41.00 Matches: 7
Percent Similarity: 100.00% Conservat: 2
Best Local Similarity: 77.78% Mismatches: 0
Query Match: 93.18% Indels: 0
DB: 28 Gaps: 0

US-09-965-594-1_COPY_14_22 (1-9) x BH241165 (1-758)

QY 1 LeuLeuGlyCysIleThrSerLeu 9
DB 130 TTGTTAGGTTGCATTGTGACGAGTATG 156

RESULT 14

LOCUS A0955329 724 bp DNA linear GSS 28-JAN-2000
DEFINITION LERAD347F LRR Arabidopsis thaliana genomic clone LERAD34, genomic survey sequence.

ACCESSION A0955329
VERSION A0955329.1 GI:6783030
KEYWORDS GSS.

SOURCE Arabidopsis thaliana (thale cress)

ORGANISM

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids

; eurosids II; Brassicales; Brassicaceae; Arabidopsis.

REFERENCE 1 (bases 1 to 724)

AUTHORS Ruel, C.R., Lin, X., Pai, G., Barnstead, M., Bowman, C., Utterback, T., Feldblyum, J., Liang, F., Creasy, J., and Fraser, C.M.

TITLE Genomic survey sequencing of Landsberg erecta ecotype of Arabidopsis thaliana and identification of sequence-based polymorphisms

JOURNAL Unpublished

COMMENT

Contact: Xiaoying Lin

The Institute for Genomic Research

9712 Medical Center Dr., Rockville, MD 20850, USA

Tel: 301 838 0200

Fax: 301 838 0208

Email: at@tigr.org

Seq primer: TF

Class: shotgun.

FEATURES

source
Location/Qualifiers

1..724
/organism="Arabidopsis thaliana"
/mol_type="genomic DNA"
/strain="Landsberg erecta"
/db_xref="taxon:3702"
/clone="LERAD34"
/clone_lib="LERA"

/note="Organ: Leaf; Vector: pPOS2; Total genomic DNA was sheared to 0.9-1 Kbp before ligation."

BASE COUNT 187 a 151 c 143 g 243 t

ORIGIN

Alignment Scores:
Pred. No.: 1,56e+03 Length: 724
Score: 41.00 Matches: 7
Percent Similarity: 100.00% Conservat: 2
Best Local Similarity: 77.78% Mismatches: 0
Query Match: 93.18% Indels: 0
DB: 28 Gaps: 0

US-09-965-594-1_COPY_14_22 (1-9) x A0955329 (1-724)

QY 1 LeuLeuGlyCysIleThrSerLeu 9

DB 70 TTGTTAGGTTGCATTGTGACGAGTATG 96

RESULT 15

LOCUS BH241165

DEFINITION BH241165

ACCESSION BH241165

VERSION BH241165.1

KEYWORDS GSS.

SOURCE Arabidopsis thaliana (thale cress)

ORGANISM Arabidopsis thaliana

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids

; eurosids II; Brassicales; Brassicaceae; Arabidopsis.

REFERENCE 1 (bases 1 to 758)

AUTHORS Town, C.D., Whitelaw, C.A., Pai, G., Van Aken, S.E., Utterback, T.V., Feldblyum, T.V. and Fraser, C.M.

TITLE Survey sequencing of Arabidopsis thaliana BAC F25J21

JOURNAL Unpublished

COMMENT Other GSSs: AULF85TR

Contact: Chris Town

TIGR

9712 Medical Center Drive, Rockville, MD 20850, USA.

Tel: 301-838-3523

Fax: 301-838-0208

Email: cdtown@tigr.org

From Wash. U contig 720.

Seq primer: TF

Class: sheared ends.

Location/Qualifiers

1..758

/organism="Arabidopsis thaliana"

/mol_type="genomic DNA"

/strain="Columbia"

/db_xref="taxon:3702"

/clone="AULF85"

/clone_lib="AULF"

/note="Vector: pPOS2; Site 1: BstXI; 2-3 kb sheared BAC DNA inserted into pPOS2 using BstXI linkers"

BASE COUNT 282 a 140 c 157 g 259 t

ORIGIN

Alignment Scores:
Pred. No.: 1,61e+03 Length: 758
Score: 41.00 Matches: 7
Percent Similarity: 100.00% Conservat: 2
Best Local Similarity: 77.78% Mismatches: 0
Query Match: 93.18% Indels: 0
DB: 28 Gaps: 0

US-09-965-594-1_COPY_14_22 (1-9) x BH241165 (1-758)

QY 1 LeuLeuGlyCysIleThrSerLeu 9

DB 318 TTGTTAGGTTGCATTGTGACGAGTATG 344

Search completed: September 29, 2003, 20:22:55
Job time : 1632 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2003 Computer Ltd.

OM protein - protein search, using sw model

Run on: September 23, 2003, 19:00:07 : Search time 30 Seconds
(without alignments)
77,416 Million cells updates/sec

Title: US-09-965-594-l_copy_14_22

Perfect score: 44

Sequence: 1 LLAGCIITSL 9

Scoring table: BLOSUM62

Gapop 13.0 , Gapext 0.5

Searched: 830525 seqs, 25902694 residues

Total number of hits satisfying chosen parameters: 830296

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 99%

Listing first 45 summaries

Database :

SPTREMBL_23:*

1: sp_archaea:*

2: sp_bacteria:*

3: sp_fungi:*

4: sp_human:*

5: sp_invertebrate:*

6: sp_mammal:*

7: sp_mmc:*

8: sp_organelle:*

9: sp_phase:*

10: sp_plant:*

11: sp_rodent:*

12: sp_virus:*

13: sp_vertebrate:*

14: sp_unclassified:*

15: sp_rvirus:*

16: sp_bacteriaph:*

17: sp_archaeap:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	43	97.7	180	12	Q8QW28
2	43	97.7	180	12	Q8QW29
3	43	97.7	180	12	Q8QW27
4	43	97.7	180	12	Q8QW23
5	43	97.7	403	12	Q8QW94
6	43	97.7	3010	12	Q9QIY5
7	43	97.7	3010	12	Q9QIY6
8	43	97.7	3010	12	Q9QIY2
9	43	97.7	3010	12	Q8V638
10	43	97.7	3010	12	Q81541
11	43	97.7	3010	12	Q9QID6
12	43	97.7	3011	12	Q91304
13	42	95.5	181	12	Q913P5
14	42	95.5	3010	12	Q9J3B9
15	42	95.5	3010	12	Q9J3G5
16	41	93.2	180	12	Q8QW19

17	41	93.2	361	12	Q70817	Q70817 hepatitis C
18	41	93.2	2864	12	Q92973	Q92973 hepatitis C
19	41	93.2	2864	12	Q9W1K8	Q9W1K8 hepatitis C
20	41	93.2	2864	12	Q92976	Q92976 hepatitis C
21	41	93.2	3010	12	Q02829	Q02829 hepatitis C
22	41	93.2	3010	12	Q92959	Q92959 hepatitis C
23	41	93.2	3010	12	Q02828	Q02828 hepatitis C
24	41	93.2	3010	12	Q9J3G0	Q9J3G0 hepatitis C
25	40	90.9	180	12	Q8QVX2	Q8QVX2 hepatitis C
26	40	90.9	180	12	Q8QW22	Q8QW22 hepatitis C
27	40	90.9	180	12	Q8QW09	Q8QW09 hepatitis C
28	40	90.9	180	12	Q8QW26	Q8QW26 hepatitis C
29	40	90.9	180	12	Q8QVW8	Q8QVW8 hepatitis C
30	40	90.9	180	12	Q8QVX0	Q8QVX0 hepatitis C
31	40	90.9	180	12	Q8QVW2	Q8QVW2 hepatitis C
32	40	90.9	180	12	Q8QV22	Q8QV22 hepatitis C
33	40	90.9	180	12	Q8QV26	Q8QV26 hepatitis C
34	40	90.9	181	12	Q91RQ7	Q91RQ7 hepatitis C
35	40	90.9	181	12	Q91RQ6	Q91RQ6 hepatitis C
36	40	90.9	2864	12	Q9W1K9	Q9W1K9 hepatitis C
37	40	90.9	2864	12	Q9W1L0	Q9W1L0 hepatitis C
38	40	90.9	2864	12	Q92975	Q92975 hepatitis C
39	40	90.9	2864	12	Q92974	Q92974 hepatitis C
40	40	90.9	3010	12	P90192	P90192 hepatitis C
41	40	90.9	3010	12	Q9C1Y7	Q9C1Y7 hepatitis C
42	40	90.9	3010	12	Q92970	Q92970 hepatitis C
43	40	90.9	3010	12	Q9J3G9	Q9J3G9 hepatitis C
44	40	90.9	3010	12	Q9J3G4	Q9J3G4 hepatitis C
45	40	90.9	3010	12	Q9QIY8	Q9QIY8 hepatitis C

ALIGNMENTS

RESULT 1

Q8QW08 ID Q8QW08 PRELIMINARY: PRI: 180 AA.
AC Q8QW08;
DI 01-JUN-2002 (TrEMBLrel. 21, Created)
DI 01-JUN-2002 (TrEMBLrel. 21, Last sequence update)
DI 01-OCT-2002 (TrEMBLrel. 22, Last annotation update)
DE Polypeptide (Fragment).
CS Hepatitis C virus type 1b.
OC Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
OC Hepacivirus.
CX NCBI_TaxID=31647;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=108;
KA Odata S.;
RT *CORRELATION BETWEEN SECONDARY STRUCTURE OF AN AMINO-TERMINAL PORTION OF THE NONSTRUCTURAL PROTEIN 3 OF HEPATITIS C VIRUS AND DEVELOPMENT OF HEPATOCELLULAR CARCINOMA.*
RL Submitted (SEP-2001) to the EMBL/GenBank/DBJ databases.
DR EMBL: AB072067; BA588249.1; -;
DR InterPro: IPR004109; HCV_NS3.
DR Pfam: PF02907; HCV_NS3.1.
FT NON_TER 1
FT NON_TER 180
SQ SEQUENCE 180 AA: 18982 MW: 77E64C2673D89EEC CRC64:

Query Match 97.7%; Score 43; DB 12; Length 180;
Best Local similarity 98.9%; Pred. No. 1.1;
Matches 8; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

OY 1 LLAGCIITSL 9
Db 13 LLAGCIITSL 21

RESULT 2

Q8QW28 ID Q8QW28 PRELIMINARY: PRI: 180 AA.

```
AC Q8QW28;
DI 01-JUN-2002 (TrEMBLrel. 21, Created)
DI 01-JUN-2002 (TrEMBLrel. 21, Last sequence update)
DI 01-OCT-2002 (TrEMBLrel. 22, Last annotation update)
DE Polyprotein (Fragment).
OS Hepatitis C virus type 1b.
OC Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
OC Hepacivirus
OX NCBI_TaxID=31647;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=H05-4;
RA Odata S.;
RT *CORRELATION BETWEEN SECONDARY STRUCTURE OF AN AMINO-TERMINAL PORTION
RT OF THE NONSTRUCTURAL PROTEIN 3 OF HEPATITIS C VIRUS AND DEVELOPMENT OF
RI HEPATOCELLULAR CARCINOMA.*;
RL Submitted (SEP-2001) to the EMBL/GenBank/DBJ databases.
DR EMBL; AB072047; BAB88229.1; -.
DR InterPro; IPR004109; HCV_NS3.
DR Pfam; PF02907; HCV_NS3; 1.
FT NON_TER 1 180
FT NON_TER 180 180
SQ SEQUENCE 180 AA: 18891 MW: 85AF5CE1CF2BCA30 CRC64;

Query Match 97.7%; Score 43; DB 12; Length 180;
Best Local Similarity 88.9%; Pred. No. 1.1;
Matches 8; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 LLGCIITSL 9
DB 13 LLGCIIVTSL 21

RESULT 3
Q8QW27
ID Q8QW27 PRELIMINARY; PRT; 180 AA.
AC Q8QW27;
DI 01-JUN-2002 (TrEMBLrel. 21, Created)
DI 01-JUN-2002 (TrEMBLrel. 21, Last sequence update)
DI 01-OCT-2002 (TrEMBLrel. 22, Last annotation update)
DE Polyprotein (Fragment).
OS Hepatitis C virus type 1b.
OC Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
OC Hepacivirus
OX NCBI_TaxID=31647;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=H05-5;
RA Odata S.;
RT *CORRELATION BETWEEN SECONDARY STRUCTURE OF AN AMINO-TERMINAL PORTION
RT OF THE NONSTRUCTURAL PROTEIN 3 OF HEPATITIS C VIRUS AND DEVELOPMENT OF
RI HEPATOCELLULAR CARCINOMA.*;
RL Submitted (SEP-2001) to the EMBL/GenBank/DBJ databases.
DR EMBL; AB072048; BAB88230.1; -.
DR InterPro; IPR004109; HCV_NS3.
DR Pfam; PF02907; HCV_NS3; 1.
FT NON_TER 1 180
FT NON_TER 180 180
SQ SEQUENCE 180 AA: 18913 MW: 895FB21253DA6CDB CRC64;

Query Match 97.7%; Score 43; DB 12; Length 180;
Best Local Similarity 88.9%; Pred. No. 1.1;
Matches 8; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 LLGCIITSL 9
DB 13 LLGCIIVTSL 21

RESULT 4
Q8QVX3
ID Q8QVX3 PRELIMINARY; PRT; 180 AA.
AC Q8QVX3;
```

```
LT 01-JUN-2002 (TrEMBLrel. 21, Created)
DI 01-JUN-2002 (TrEMBLrel. 21, Last sequence update)
DI 01-OCT-2002 (TrEMBLrel. 22, Last annotation update)
DE Polyprotein (Fragment).
OS Hepatitis C virus type 1b.
OC Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
OC Hepacivirus
OX NCBI_TaxID=31647;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=N-82;
RA Odata S.;
RT *CORRELATION BETWEEN SECONDARY STRUCTURE OF AN AMINO-TERMINAL PORTION
RT OF THE NONSTRUCTURAL PROTEIN 3 OF HEPATITIS C VIRUS AND DEVELOPMENT OF
RI HEPATOCELLULAR CARCINOMA.*;
RL Submitted (SEP-2001) to the EMBL/GenBank/DBJ databases.
DR EMBL; AB072102; BAB88284.1; -.
DR InterPro; IPR004109; HCV_NS3.
DR Pfam; PF02907; HCV_NS3; 1.
FT NON_TER 1 180
FT NON_TER 180 180
SQ SEQUENCE 180 AA: 18808 MW: CE520CF0BCA4E1E2 CRC64;

Query Match 97.7%; Score 43; DB 12; Length 180;
Best Local Similarity 88.9%; Pred. No. 1.1;
Matches 8; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 LLGCIITSL 9
DB 13 LLGCIIVTSL 21

RESULT 5
Q8QF94
ID Q8QF94 PRELIMINARY; PRT; 403 AA.
AC Q8QF94;
DI 01-JUN-2002 (TrEMBLrel. 21, Created)
DI 01-JUN-2002 (TrEMBLrel. 21, Last sequence update)
DI 01-MAR-2003 (TrEMBLrel. 23, Last annotation update)
DE Genome Polyprotein (Fragment).
OS Hepatitis C virus.
OC Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
OC Hepacivirus
OX NCBI_TaxID=11103;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=667;
RA MEELINE-21904745; PubMed=1197242;
RA Kallings O., Norder H., Mukomolov S., Magnus L.O.;
RT *A natural intergenotypic recombinant of hepatitis C virus identified
RT in St. Petersburg.*;
PL J. Virol. 76:4034-4043(2002).
DR EMBL; AY070171; AAL58587.1; -.
DR InterPro; IPR02531; HCV_NS1.
DR InterPro; IPR02518; HCV_NS2.
DR InterPro; IPR004109; HCV_NS3.
DR Pfam; PF01560; HCV_NS1; 1.
DR Pfam; PF01538; HCV_NS2; 1.
DR Pfam; PF02907; HCV_NS3; 1.
KW Coat protein; Envelope protein; Glycoprotein; Nonstructural protein;
KW Polyprotein; Transmembrane.
FT NON_TER 1 403
FT NON_TER 403 403
SQ SEQUENCE 403 AA: 43868 MW: EASCFACB9348C8A CRC64;

Query Match 97.7%; Score 43; DB 12; Length 403;
Best Local Similarity 88.9%; Pred. No. 2.2;
Matches 8; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 LLGCIITSL 9
DB 323 LLGCIIVTSL 331
```

RESULT 6

Q90IY5 PRELIMINARY: PRT: 3010 AA.
 AC Q90IY5: (1)
 DT 01-MAY-2000 (TRENBLREL. 23, Created)
 DI 01-MAY-2000 (TRENBLREL. 23, Last sequence update)
 DE Genome polyprotein.
 OS Hepatitis C virus.
 OC Viruses: ssRNA positive-strand viruses, no DNA stage: Flaviviridae;
 OC Hepacivirus.
 OX NCBI_TaxID=11103;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=MD3-2;
 RX MEDLINE=20013325; PubMed=12544098;
 RA Nagayama K., Kurosaki M., Enomoto N., Maekawa S.Y., Miyasaka Y.,
 RA Tazawa J.I., Izumi N., Marumo F., Sato C.;
 RT "Time-related changes in full-length hepatitis C virus and hepatitis
 activity.";
 RL Virology 263:244-253(1999).
 RN [2]
 RP SEQUENCE FROM N.A.
 RC STRAIN=MD3-2;
 RA Nagayama K., Kurosaki M., Enomoto N., Maekawa S., Miyasaka Y.,
 RA Sakamoto N., Fukuma T., Tazawa J., Izumi N., Marumo F., Sato C.;
 RL Submitted (JUL-1999) to the EMBL/GenBank/DBJ databases.
 CC -1- SUBUNIT: THE VIRION OF THIS VIRUS IS A NUCLEOCAPSID COVERED BY A
 CC LIPID PROTEIN ENVELOPE. THE ENVELOPE CONSISTS OF TWO PROTEINS:
 CC PROTEIN M AND GLYCOPROTEIN E. THE NUCLEOCAPSID IS A COMPLEX OF
 CC PROTEIN C AND MRNA (BY SIMILARITY).
 CC EMBL: AF155050; AAD56185.1; .
 DR HSP: P26563; IJXP.
 DR InterPro: IPR001410; DEAD.
 DR InterPro: IPR002522; HCV_capsid.
 DR InterPro: IPR002521; HCV_core.
 DR InterPro: IPR002519; HCV_env.
 DR InterPro: IPR002531; HCV_NS1.
 DR InterPro: IPR002518; HCV_NS2.
 DR InterPro: IPR004109; HCV_NS3.
 DR InterPro: IPR000745; HCV_NS4a.
 DR InterPro: IPR001490; HCV_NS4b.
 DR InterPro: IPR002868; HCV_NS5a.
 DR InterPro: IPR002166; HCV_RDRP.
 DR InterPro: IPR007095; RNA_pol_PS.
 DR InterPro: IPR007034; RNA_pol_PSVir.
 DR Pfam: PF01543; HCV_capsid; 1.
 DR Pfam: PF01542; HCV_core; 1.
 DR Pfam: PF01539; HCV_env; 1.
 DR Pfam: PF01560; HCV_NS1; 1.
 DR Pfam: PF01538; HCV_NS2; 1.
 DR Pfam: PF02907; HCV_NS3; 1.
 DR Pfam: PF01066; HCV_NS4a; 1.
 DR Pfam: PF01001; HCV_NS4b; 1.
 DR Pfam: PF01506; HCV_NS5a; 1.
 DR Pfam: PF00998; Viral_RDRP; 1.
 DR ProDom: PD186062; HCV_NS1; 1.
 DR SMART: SM00487; DEXDC; 1.
 DR PROSITE: PS05057; RDRP_POSITIVE; 1.
 DR PROSITE: PS05052; RDRP_VIRAL; 1.
 KW Coat protein: Envelope protein: Glycoprotein: Nonstructural protein:
 KW Polyprotein: RNA-directed RNA polymerase: Transferase: Transmembrane.
 SQ SEQUENCE 3010 AA: 327430 MW: 15190E3463DRAC15 CRC64;

Query Match 97.7% Score 43; DB 12; Length 3010;
 Best Local Similarity 88.9%; Pred. No. 12;
 Matches 8; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 LLGCIVTSL 9

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Db 1039 LLGCIVTSL 1047

RESULT 7

Q90IY6 PRELIMINARY: PRT: 3010 AA.
 AC Q90IY6: (1)
 DT 01-MAY-2000 (TRENBLREL. 13, Created)
 DI 01-MAY-2000 (TRENBLREL. 13, Last sequence update)
 DE Genome polyprotein.
 OS Hepatitis C virus.
 OC Viruses: ssRNA positive-strand viruses, no DNA stage: Flaviviridae;
 OC Hepacivirus.
 OX NCBI_TaxID=11103;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=MD3-1;
 RX MEDLINE=20013325; PubMed=10544098;
 RA Nagayama K., Kurosaki M., Enomoto N., Maekawa S.Y., Miyasaka Y.,
 RA Tazawa J.I., Izumi N., Marumo F., Sato C.;
 RT "Time-related changes in full-length hepatitis C virus and hepatitis
 activity.";
 RL Virology 263:244-253(1999).
 RN [2]
 RP SEQUENCE FROM N.A.
 RC STRAIN=MD3-1;
 RA Nagayama K., Kurosaki M., Enomoto N., Maekawa S., Miyasaka Y.,
 RA Sakamoto N., Fukuma T., Tazawa J., Izumi N., Marumo F., Sato C.;
 RL Submitted (JUL-1999) to the EMBL/GenBank/DBJ databases.
 CC -1- SUBUNIT: THE VIRION OF THIS VIRUS IS A NUCLEOCAPSID COVERED BY A
 CC LIPID PROTEIN ENVELOPE. THE ENVELOPE CONSISTS OF TWO PROTEINS:
 CC PROTEIN M AND GLYCOPROTEIN E. THE NUCLEOCAPSID IS A COMPLEX OF
 CC PROTEIN C AND MRNA (BY SIMILARITY).
 CC EMBL: AF155049; AAD56184.1; .
 DR HSP: P26663; IJXP.
 DR InterPro: IPR001410; DEAD.
 DR InterPro: IPR002522; HCV_capsid.
 DR InterPro: IPR002521; HCV_core.
 DR InterPro: IPR002519; HCV_env.
 DR InterPro: IPR002531; HCV_NS1.
 DR InterPro: IPR002518; HCV_NS2.
 DR InterPro: IPR004109; HCV_NS3.
 DR InterPro: IPR000745; HCV_NS4a.
 DR InterPro: IPR001490; HCV_NS4b.
 DR InterPro: IPR002868; HCV_NS5a.
 DR InterPro: IPR002166; HCV_RDRP.
 DR InterPro: IPR007095; RNA_pol_DS_PS.
 DR InterPro: IPR007034; RNA_pol_PSVir.
 DR Pfam: PF01543; HCV_capsid; 1.
 DR Pfam: PF01542; HCV_core; 1.
 DR Pfam: PF01539; HCV_env; 1.
 DR Pfam: PF01560; HCV_NS1; 1.
 DR Pfam: PF01538; HCV_NS2; 1.
 DR Pfam: PF02907; HCV_NS3; 1.
 DR Pfam: PF01066; HCV_NS4a; 1.
 DR Pfam: PF01001; HCV_NS4b; 1.
 DR Pfam: PF01506; HCV_NS5a; 1.
 DR Pfam: PF00998; Viral_RDRP; 1.
 DR ProDom: PD186062; HCV_NS1; 1.
 DR SMART: SM00487; DEXDC; 1.
 DR PROSITE: PS05057; RDRP_POSITIVE; 1.
 DR PROSITE: PS05052; RDRP_VIRAL; 1.
 KW Coat protein: Envelope protein: Glycoprotein: Nonstructural protein:
 KW Polyprotein: RNA-directed RNA polymerase: Transferase: Transmembrane.
 SQ SEQUENCE 3010 AA: 327368 MW: 998C7F293EAAEC8D CRC64;

Query Match 97.7% Score 43; DB 12; Length 3010;
 Best Local Similarity 88.9%; Pred. No. 12;
 Matches 8; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 LLGCIVTSL 9

:|||||:

Db 1039 LLGCIVTSL 1047

RESULT 8

Q9QIY2 PRELIMINARY: PRT: 3010 AA.
 AC Q9QIY2
 DT 01-MAY-2000 (TrEMBLrel. 13, Created)
 DT 01-MAY-2000 (TrEMBLrel. 13, Last sequence update)
 DI 01-MAR-2003 (TrEMBLrel. 23, Last annotation update)
 DE Genome polyprotein.
 OS Hepatitis C virus.
 OC Viruses; ssRNA positive-strand viruses, no RNA stage; Flaviviridae;
 OC Hepacivirus.
 OX NCBI_TaxID=11103;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=MD5-1;
 RX MEDLINE=20013325; PubMed=10544098;
 RA Nagayama K., Kurosaki M., Enomoto N., Maekawa S., Miyasaka Y.,
 RA Tazawa J.-i., Izumi N., Marumo F., Sato C.;
 RT "Time-related changes in full-length hepatitis C virus and hepatitis
 RT activity";
 RL Virology 263:244-253 (1999).
 RN [2]
 RP SEQUENCE FROM N.A.
 RC STRAIN=MD5-1;
 RA Nagayama K., Kurosaki M., Enomoto N., Maekawa S., Miyasaka Y.,
 RA Sakamoto N., Fukuma I., Tazawa J., Izumi N., Marumo F., Sato C.;
 RL Submitted (JUL-1999) to the EMBL/GenBank/DBJ databases.
 CC -1- SUBUNIT: THE VIRION OF THIS VIRUS IS A NUCLEOCAPSID COVERED BY A
 CC LIPOPROTEIN ENVELOPE. THE ENVELOPE CONSISTS OF TWO PROTEINS:
 CC PROTEIN M AND GLYCOPROTEIN E. THE NUCLEOCAPSID IS A COMPLEX OF
 CC PROTEIN C AND MRNA (BY SIMILARITY).
 DR EMBL: AF165053; AAD56188.1;
 DR HSP: P27958; IHEI.
 DR InterPro: IPR001410; DEAD.
 DR InterPro: IPR002522; HCV_capsid.
 DR InterPro: IPR002521; HCV_core.
 DR InterPro: IPR002531; HCV_env.
 DR InterPro: IPR002518; HCV_NS1.
 DR InterPro: IPR004100; HCV_NS2.
 DR InterPro: IPR001490; HCV_NS4a.
 DR InterPro: IPR002868; HCV_NS5a.
 DR InterPro: IPR002166; HCV_RdRp.
 DR InterPro: IPR007095; RNA_pol_DS_PS.
 DR InterPro: IPR007094; RNA_pol_PSVir.
 DR Pfam: PF01543; HCV_capsid; 1.
 DR Pfam: PF01542; HCV_core; 1.
 DR Pfam: PF01539; HCV_env; 1.
 DR Pfam: PF01565; HCV_NS1; 1.
 DR Pfam: PF01533; HCV_NS2; 1.
 DR Pfam: PF02907; HCV_NS3; 1.
 DR Pfam: PF01006; HCV_NS4a; 1.
 DR Pfam: PF01001; HCV_NS4b; 1.
 DR Pfam: PF01506; HCV_NS5a; 1.
 DR Pfam: PF00998; Viral_RdRp; 1.
 DR ProDom: PD186062; HCV_NS1; 1.
 DR Pfam: PF01539; HCV_env; 1.
 DR Pfam: PF01565; HCV_NS1; 1.
 DR Pfam: PF01538; HCV_NS2; 1.
 DR Pfam: PF02907; HCV_NS3; 1.
 DR Pfam: PF01006; HCV_NS4a; 1.
 DR Pfam: PF01001; HCV_NS4b; 1.
 DR Pfam: PF01506; HCV_NS5a; 1.
 DR Pfam: PF00998; Viral_RdRp; 1.
 DR ProDom: PD186062; HCV_NS1; 1.
 DR SMART: SM00487; DEXDC; 1.
 DR PROSITE: PS00507; RDRP_POSITIVE; 1.
 DR PROSITE: PS05021; RDRP_VIRAL; 1.
 KW Coat protein; Envelope protein; Glycoprotein; Nonstructural protein;
 KW Polyprotein; RNA-directed RNA polymerase; Transferase; Transmembrane.
 FT NON_TER 3010
 SQ SEQUENCE 3010 AA; 326595 MW; D186BA7E2F0B5E8 CUC64;
 Query Match 97.7%; Score 43; DB 12; Length 3010;
 Best Local Similarity 88.9%; Pred. No. 12;
 Matches 8; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

Qy 1 LLGCITSL 9

Db 1039 LLGCITSL 1047
 |||||:||||

RESULT 9

Q8V638 PRELIMINARY: PRT: 3010 AA.
 AC Q8V638;
 DT 01-MAR-2002 (TrEMBLrel. 20, Created)
 DT 01-MAR-2002 (TrEMBLrel. 20, Last sequence update)
 DI 01-MAR-2003 (TrEMBLrel. 23, Last annotation update)
 DE Genome polyprotein (Fragment).
 OS Hepatitis C virus.
 OC Viruses; ssRNA positive-strand viruses, no RNA stage; Flaviviridae;
 OC Hepacivirus.
 OX NCBI_TaxID=11103;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC Fanning L.J., Itakura J., Nagayama K., Enomoto N.;
 RT "Characteristics of Hepatitis C viral genome associated with disease
 RT progression in a homogeneous patient population";
 RL Submitted (OCT-2000) to the EMBL/GenBank/DBJ databases.
 CC -1- SUBUNIT: THE VIRION OF THIS VIRUS IS A NUCLEOCAPSID COVERED BY A
 CC LIPOPROTEIN ENVELOPE. THE ENVELOPE CONSISTS OF TWO PROTEINS:
 CC PROTEIN M AND GLYCOPROTEIN E. THE NUCLEOCAPSID IS A COMPLEX OF
 CC PROTEIN C AND MRNA (BY SIMILARITY).
 DR EMBL: AF313916; AAL55821.1;
 DR InterPro: IPR000345; CytC_heme_bind.
 DR InterPro: IPR00410; DEAD.
 DR InterPro: IPR002522; HCV_capsid.
 DR InterPro: IPR002521; HCV_core.
 DR InterPro: IPR002519; HCV_env.
 DR InterPro: IPR002533; HCV_NS1.
 DR InterPro: IPR002518; HCV_NS2.
 DR InterPro: IPR004109; HCV_NS3.
 DR InterPro: IPR003745; HCV_NS4a.
 DR InterPro: IPR001490; HCV_NS4b.
 DR InterPro: IPR002868; HCV_NS5a.
 DR InterPro: IPR002166; HCV_RdRp.
 DR InterPro: IPR007095; RNA_pol_DS_PS.
 DR InterPro: IPR007094; RNA_pol_PSVir.
 DR Pfam: PF01543; HCV_capsid; 1.
 DR Pfam: PF01542; HCV_core; 1.
 DR Pfam: PF01539; HCV_env; 1.
 DR Pfam: PF01565; HCV_NS1; 1.
 DR Pfam: PF01533; HCV_NS2; 1.
 DR Pfam: PF02907; HCV_NS3; 1.
 DR Pfam: PF01006; HCV_NS4a; 1.
 DR Pfam: PF01001; HCV_NS4b; 1.
 DR Pfam: PF01506; HCV_NS5a; 1.
 DR Pfam: PF00998; Viral_RdRp; 1.
 DR ProDom: PD186062; HCV_NS1; 1.
 DR SMART: SM00487; DEXDC; 1.
 DR PROSITE: PS00190; CYTOCHROME_C; 1.
 DR PROSITE: PS05007; RDRP_POSITIVE; 1.
 DR PROSITE: PS05021; RDRP_VIRAL; 1.
 KW Coat protein; Envelope protein; Glycoprotein; Nonstructural protein;
 KW Polyprotein; RNA-directed RNA polymerase; Transferase; Transmembrane.
 FT NON_TER 3010
 SQ SEQUENCE 3010 AA; 327181 MW; 33AAA6C07251C839 CRC64;
 Query Match 97.7%; Score 43; DB 12; Length 3010;
 Best Local Similarity 88.9%; Pred. No. 12;
 Matches 8; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

Qy 1 LLGCITSL 9
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 Db 1039 LLGCITSL 1047

RESULT 10

Q81541

ID Q81541; PRELIMINARY: PRT: 3010 AA.

AC Q81541;

DT 01-NOV-1996 (TrEMBLrel. 01, Created)

DT 01-NOV-1996 (TrEMBLrel. 01, Last sequence update)


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DT 01-DEC-2001 (TrEMBLrel. 19, Created;
DT 01-DEC-2001 (TrEMBLrel. 19, Last sequence update)
DT 01-MAR-2003 (TrEMBLrel. 23, Last annotation update)
DE Genome polyprotein.
OS Hepatitis C virus.
OC Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
OC Hepacivirus.
OX NCBI_TaxID=11103;
RN [1]
RP SEQUENCE FROM N.A.
RA Guntaka K.V., Kumpally S.K., Khaja M.N., Kota K.K., Ramana V.K.,
RA Swaminathan S., Sakata Y., Habeebullah C.M.;
RT "Nucleotide sequence of Indian strain of Hepatitis C Virus.";
RL Submitted (AUG-2001) to the EMBL/GenBank/DBJ databases.
CC -1- SUBUNIT: THE VIRION OF THIS VIRUS IS A NUCLEOCAPSID COVERED BY A
CC LIPOPROTEIN ENVELOPE. THE ENVELOPE CONSISTS OF TWO PROTEINS:
CC PROTEIN M AND GLYCOPROTEIN E. THE NUCLEOCAPSID IS A COMPLEX OF
CC PROTEIN C AND MRNA (BY SIMILARITY).
DR EMBL: AY051292; AAK95932.1;
DR MEROPS: S29.001;
DR MEROPS: U39.001;
DR InterPro: IPR000345; CytC_heme_bind.
DR InterPro: IPR001410; DEAD.
DR InterPro: IPR002522; HCV_capsid.
DR InterPro: IPR002521; HCV_core.
DR InterPro: IPR002519; HCV_env.
DR InterPro: IPR002531; HCV_NS1.
DR InterPro: IPR002518; HCV_NS2.
DR InterPro: IPR004109; HCV_NS3.
DR InterPro: IPR000745; HCV_NS4a.
DR InterPro: IPR001490; HCV_NS4b.
DR InterPro: IPR002868; HCV_NS5a.
DR InterPro: IPR032166; HCV_RdRP.
DR InterPro: IPR007095; RNA_pol_DS_PS.
DR InterPro: IPR037694; RNA_pol_PSVir.
DR Pfam: PF01543; HCV_capsid; 1.
DR Pfam: PF01542; HCV_core; 1.
DR Pfam: PF01560; HCV_NS1; 1.
DR Pfam: PF02907; HCV_NS3; 1.
DR Pfam: PF01006; HCV_NS4a; 1.
DR Pfam: PF01001; HCV_NS4b; 1.
DR Pfam: PF01506; HCV_NS5a; 1.
DR Pfam: PF00398; Viral_RdRP; 1.
DR ProDom: PD186062; HCV_NS1; 1.
DR SMART: SM00487; DEXDC; 1.
DR PROSITE: PS00190; CYTOCHROME_C; 1.
DR PROSITE: PS0507; RDRP_POSITIVE; 1.
DR PROSITE: PS0521; RDRP_VIRAL; 1.
KW Coat protein: envelope protein; Glycoprotein: Nonstructural protein.
KW Polyprotein: RNA-directed RNA polymerase; Transcriptase; Transmembrane.
SQ SEQUENCE 3011 AA: 327234 MW: 57A21964a422785C CRC64;
```

Query Match 97.7%; Score 43; DB 12; Length 3011;
Best Local Similarity 88.9%; Pred. No. 12;
Matches 8; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

OY 1 LLGCIITSL 9
DB 1039 LLGCIIVTSL 1047

RESULT 13

OY1RP5 Q91RP5 PRELIMINARY: PRT; 181 AA.

ID Q91RP5
AC Q91RP5
DT 01-DEC-2001 (TrEMBLrel. 19, Created)
DT 01-DEC-2001 (TrEMBLrel. 19, Last sequence update)
DT 01-MAR-2003 (TrEMBLrel. 23, Last annotation update)
DE NS3 protease (fragment).
OS Hepatitis C virus.
OC Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;

```
OC Hepacivirus.
OX NCBI_TaxID=11103;
RN [1]
RP SEQUENCE FROM N.A.
RA Holland-Staley C.A., Kovari L.C., Golenberg E., Meyers D.L.;
RT "Genetic Diversity and Response to IFN of the NS3 Protease Gene from
RT Clinical Strains of the Hepatitis C Virus.";
RL Submitted (APR-2001) to the EMBL/GenBank/DBJ databases.
DR EMBL: AF369258; AAK54583.1;
DR MEROPS: S29.001;
DR MEROPS: U39.001;
DR InterPro: IPR004109; HCV_NS3.
DR Pfam: PF02907; HCV_NS3; 1.
KW Protease.
PT NON_TER 1 181
FT NON_TER 181 181
SQ SEQUENCE 181 AA: 19056 MW: BAF89690AD2971DB CRC64;

Query Match 95.5%; Score 42; DB 12; Length 181;  
Best Local Similarity 88.9%; Pred. No. 1.8;  
Matches 8; Conservative 1; Mismatches 0; Indels 0; Gaps 0;



OY 1 LLGCIITSL 9  
DB 13 ILGCIITSL 21



RESULT 14



OY13H5 Q9J3H5 PRELIMINARY: PRT; 3010 AA.



ID Q9J3H5  
AC Q9J3H5  
DT 01-OCT-2000 (TrEMBLrel. 15, Created)  
DT 01-OCT-2000 (TrEMBLrel. 15, Last sequence update)  
DT 01-MAR-2003 (TrEMBLrel. 23, Last annotation update)  
DE Genome polyprotein.  
OS Hepatitis C virus.  
OC Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;  
OC Hepacivirus.  
OX NCBI_TaxID=11103;
AN [1]
RP SEQUENCE FROM N.A.
RA Nagayama K., Kurosaki M., Enomoto N., Miyasaka Y., Marume F., Sato C.;
RT "Characteristics of hepatitis C virus genome associated with disease
RT progression.";
RL Submitted (NOV-1999) to the EMBL/GenBank/DBJ databases.
CC -1- SUBUNIT: THE VIRION OF THIS VIRUS IS A NUCLEOCAPSID COVERED BY A
CC LIPOPROTEIN ENVELOPE. THE ENVELOPE CONSISTS OF TWO PROTEINS:
CC PROTEIN M AND GLYCOPROTEIN E. THE NUCLEOCAPSID IS A COMPLEX OF
CC PROTEIN C AND MRNA (BY SIMILARITY).
DR EMBL: AF207734; AAF65944.1;
DR HSSP: P26663; IUXP.
DR InterPro: IPR000345; CytC_heme_bind.
DR InterPro: IPR001410; DEAD.
DR InterPro: IPR002522; HCV_capsid.
DR InterPro: IPR002521; HCV_core.
DR InterPro: IPR002519; HCV_env.
DR InterPro: IPR002531; HCV_NS1.
DR InterPro: IPR002518; HCV_NS2.
DR InterPro: IPR004109; HCV_NS3.
DR InterPro: IPR000745; HCV_NS4a.
DR InterPro: IPR001490; HCV_NS4b.
DR InterPro: IPR002868; HCV_NS5a.
DR InterPro: IPR002156; HCV_RdRP.
DR InterPro: IPR007095; RNA_pol_DS_PS.
DR InterPro: IPR007094; RNA_pol_PSVir.
DR Pfam: PF01543; HCV_capsid; 1.
DR Pfam: PF01542; HCV_core; 1.
DR Pfam: PF01539; HCV_env; 1.
DR Pfam: PF01560; HCV_NS1; 1.
DR Pfam: PF01538; HCV_NS2; 1.
DR Pfam: PF02907; HCV_NS3; 1.


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DR Pfam: PF01006; HCV_NS4a; 1.
DR Pfam: PF01001; HCV_NS4b; 1.
DR Pfam: PF01506; HCV_NS5a; 1.
DR Pfam: PF00998; Viral_RdRp; 1.
DR ProDom: PD186062; HCV_NS1; 1.
DR SMART: SM00487; DEXDC; 1.
DR PROSITE: PS00190; CYTOCHROME_C; 1.
DR PROSITE: PS05050; RDRP_POSITIVE; 1.
DR PROSITE: PS05052; RDRP_VIRAL; 1.
DR ProDom: PD186062; DEXDC; 1.
DR SMART: SM00487; DEXDC; 1.
DR PROSITE: PS00190; CYTOCHROME_C; 1.
DR PROSITE: PS05050; RDRP_POSITIVE; 1.
DR PROSITE: PS05052; RDRP_VIRAL; 1.
KW Coat protein; Envelope protein; Glycoprotein; Nonstructural protein;
KW Polyprotein; RNA-directed RNA polymerase; Transferase; Transmembrane.
SC SEQUENCE 3010 AA; 326984 MW; AF120606048078 CR664;

Query Match 95.5%; Score 42; DB 12; Length 3010;
Best Local Similarity 88.9%; Pred. No. 19;
Matches 8; Conservativity 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 LIGCITSL 9
DB 1039 MGCITSL 1047

RESULT 15
Q9J3G5 PRELIMINARY: PRG: 3010 AA.
AC Q9J3G5;
DT 01-OCT-2003 (Tremblere, 15, Created)
DI 01-OCT-2003 (Tremblere, 15, Last sequence update)
DI 01-MAR-2003 (Tremblere, 23, Last annotation update)
DE Genome polyprotein.
OS Hepatitis C virus.
OC Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
OC Hepacivirus.
OX NCBI_TaxID:11103;
RN [1]
RC STRAIN:MD27;
RT Nagayama K., Kurosaki M., Enomoto N., Miyasaka Y., Marumo F., Sato C.;
RT *Characteristics of hepatitis C viral genome associated with disease
RT progression*.
RL Submitted (NOV-1999) to the EMBL/GenBank/DBJ databases.
CC -!- SUBUNIT: THE VIRION OF THIS VIRUS IS A NUCLEOCAPSID COVERED BY A
CC LIPOPROTEIN ENVELOPE. THE ENVELOPE CONSISTS OF TWO PROTEINS:
CC PROTEIN M AND GLYCOPROTEIN E. THE NUCLEOCAPSID IS A COMPLEX OF
CC PROTEIN C AND MRNA (BY SIMILARITY).
DR EMBL: AF207768; AAF65958.1;
DR HSP; P26663; IJXP;
DR InterPro: IPR000345; CytC_hemo_bind.
DR InterPro: IPR001410; DEAD.
DR InterPro: IPR002522; HCV_capsid.
DR InterPro: IPR002521; HCV_core.
DR InterPro: IPR002519; HCV_env.
DR InterPro: IPR002531; HCV_NS1.
DR InterPro: IPR002518; HCV_NS2.
DR InterPro: IPR004109; HCV_NS3.
DR InterPro: IPR000745; HCV_NS4a.
DR InterPro: IPR001490; HCV_NS4b.
DR InterPro: IPR002868; HCV_NS5a.
DR InterPro: IPR002166; HCV_RdRp.
DR InterPro: IPR001650; Helicase_C.
DR InterPro: IPR007095; RNA_pol_PS_PS.
DR InterPro: IPR007294; RNA_pol_PSVit.
DR Pfam: PF01543; HCV_capsid; 1.
DR Pfam: PF01542; HCV_core; 1.
DR Pfam: PF01539; HCV_env; 1.
DR Pfam: PF01560; HCV_NS1; 1.
DR Pfam: PF01538; HCV_NS2; 1.
DR Pfam: PF02907; HCV_NS3; 1.
DR Pfam: PF01006; HCV_NS4a; 1.
DR Pfam: PF01001; HCV_NS4b; 1.
DR Pfam: PF01506; HCV_NS5a; 1.
DR Pfam: PF00271; Helicase_C; 1.
DR Pfam: PF00998; Viral_RdRp; 1.

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DR ProDom: PD186062; HCV_NS1; 1.
DR SMART: SM00487; DEXDC; 1.
DR PROSITE: PS00190; CYTOCHROME_C; 2.
DR PROSITE: PS05050; RDRP_POSITIVE; 1.
DR PROSITE: PS05052; RDRP_VIRAL; 1.
DR KW ATP-binding; Coat protein; Envelope protein; Glycoprotein; Helicase;
KW Hydrolase; Nonstructural protein; Polyprotein;
KW RNA-directed RNA polymerase; Transferase; Transmembrane.
SC SEQUENCE 3010 AA; 327777 MW; 47A51DD7678DE62F CR664;

Query Match 95.5%; Score 42; DB 12; Length 3010;
Best Local Similarity 88.9%; Pred. No. 19;
Matches 8; Conservativity 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 LIGCITSL 9
DB 1039 MGCITSL 1047

Search completed: September 29, 2003, 19:08:26
Job time : 32 secs

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